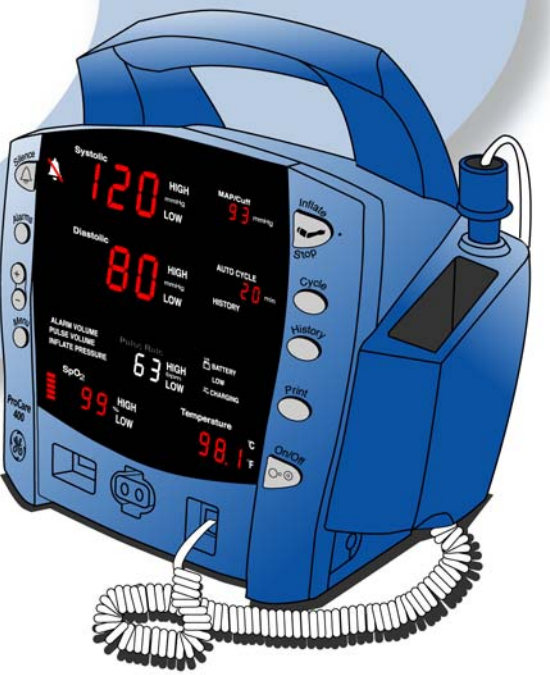
GE Medical Systems
Information Technologies

We bring good things to life.

DINAMAP®
ProCare Monitor
Operation Manual



DINAMAP®
ProCare Monitor
Operation Manual

GE Medical Systems
Information Technologies

We bring good things to life.
gemedicalsystems.com

Contents

Introduction

Reissues and Updates 3
Errors and Omissions 3
Hierarchy of Warnings and Cautions 4
About the DINAMAP® *ProCare* Monitor 7
Product Compliance 10
Symbols 11

Getting Started

Unpacking the Monitor and Accessories 15
Setting up BP Connections 15
Setting up SpO_2 Connections 16
Setting up Temperature Connections 16
Setting up the Printer (Installing the Paper) 17
Power Sources 17
Powering the Monitor: Battery and DC Supply 17
Replacing the Battery 19
Switching the Monitor On and Off 20
Setting the Date and Time 20

Product Overview

Buttons 25
Front Panel 27
Rear Panel 29
Left-Side Panel 30
Right-Side Panel 30
Windows 31
Indicators 31
Operating Modes 31
    Entering Configuration Mode 31
Parameter Modes 32
User Modes 32
Sounds 33
Power Sources 34
Specifications 35
    Mechanical 35
    Power Requirements 35
    Environmental 36

Printer

Description 41
Installing the Paper 41
Print Button 41
Printer Alarms 42
Storage 42

Alarms

Alarm Codes 45
Adjusting Alarm Limits 45
    Alarms Button 45

Adjusting the Alarm Volume 46
Silencing and Acknowledging an Alarm 46
Silence Button 46
Alarm Sounds 46
Alarm Detection and Priorities 47
Limit Alarms 47
Parameter Status Alarms 48
Printer Alarms 48
Battery Alarms 48
Memory Alarms 48
System Failure Alarms 48
Specifications 49
Alarm Message Codes 50

History

Description 55
Buttons Associated with History 55
Erasing Stored History 55
Windows Associated with History 56
Indicators Associated with History 56

BP

Description 59
Buttons Associated with BP 62
Windows Associated with BP 62
Indicators Associated with BP 63
Parameter Modes 63
BP Modes of Operation 64
Manual BP Determinations 64
Auto Cycle Determinations 64
STAT BP Determinations 65
User Settings 66
Mode Settings 66
Limit Settings 66
Neonate Settings 66
Menu Settings 67
Sounds Associated with B 67
Procedures 67
Cuff Connections 69
Quick-Disconnect Air Hose 69
Neonate Air Hose 69
Alarms 70
Critikon US Patents 70
European US Patents 70
Specifications 71
Factory Default Settings 71

SpO_2

NELLCOR® SpO_2 74
Description 74

Contents

Configuration Settings Associated with SpO_2 76
Buttons Associated with SpO_2 77
Windows Associated with SpO_2 77
Indicators Associated with SpO_2 77
Parameter Modes 77
User Settings 78
Limit settings 78
Menu Settings 78
Sounds Associated with SpO_2 78
Procedures 78
Alarms 80
Troubleshooting 81
Specifications 84
Measurement Range 84
Accuracy and Motion Tolerance 84
Saturation 84
Pulse Rate 84
NELLCOR® Sensor Accuracy 84
OxiMAX 84
OxiCliq 84
Reusable Sensor Models 85
Factory Default Settings 85
NELLCOR® Patents 85
Masimo Set® SpO_2 87
Description 87
Configuration Settings Associated with SpO_2 90
Buttons Associated with SpO_2 91
Windows Associated with SpO_2 91
Indicators Associated with SpO_2 91
Parameter Modes 91
User Settings 92
Limit settings 92
Menu Settings 92
Sounds Associated with SpO_2 92
Procedures 92
Alarms 94
Troubleshooting 95
Specifications 98
Measurement Range 98
Accuracy and Motion Tolerance 98
Saturation 98
Pulse Rate 98
Masimo® Sensor Accuracy 99
Resolution 99
Low Perfusion Performance 99
Interfering Substances 99
Sensor Light Source 99
Factory Default Settings 100
Masimo® Patents 100

TURBO★TEMP™

Description 103
Predictive Mode 103
Monitor Mode 104
Configuration Settings Associated with Temperature 105
Buttons Associated with Temperature 105
Windows Associated with Temperature 105
Indicators Associated with Temperature 105
Measurement in Progress Indicators 105
Predictive Mode 105
Measurement NOT in Progress Indicators 106
Parameter Modes 106
User Settings 107
Menu Settings 107
Sounds Associated with TURBO´TEMP™ 107
Procedures for Oral Predictive Mode Determinations 107
Procedures for Rectal Predictive Mode Determinations 108
Procedures for Monitor Mode Determinations (Axillary Determinations) 109
Alarms 111
Specifications 113
Factory Default Settings 113
IVAC® Patents 113

Pulse Rate

Description 117
General Notes 117
Buttons Associated with Pulse Rate 117
Windows Associated with Pulse Rate 118
Indicators Associated with Pulse Rate 118
Parameter Modes 118
User Settings 118
Limit settings 118
Menu Settings 118
Sounds Associated with Pulse Rate 118
Alarms 119
Specifications 120
Factory Default Settings 120

Appendix A: Principles of Noninvasive Blood Pressure

Principles of Noninvasive Blood Pressure Determination 123
Systolic Search 124
Reference Used to Determine NIBP Accuracy 125
DINAMAP Monitors With Intra-Arterial Reference (DINAMAP Classic Technology) 125
DINAMAP Monitors With Auscultatory Reference (DINAMAP Auscultatory Technology) 125

Appendix B: Maintenance

Maintenance 129
Cleaning the Monitor 129

Contents

Cuff Cleaning and Disinfection 129
General 129
Materials 129
Procedure 130
Temperature Devices 130
SpO_2 Sensors 130
Storage and Battery Care 131
Fuses 131
Calibration 131
Leak Testing 131
Disposal of Product Waste 133
Batteries 133
Patient Applied Parts 133
Packaging Material 133

Appendix C: Connection Details

Connection Details 137
Host Port Connector (rear panel) 137

Appendix D: Warranty, Service, & Spare Parts

Warranty, Service, and Spare Parts 141
Warranty 141
Assistance and Parts 141
Repairs 142
Packing Instructions 142
Service Manuals 142

Appendix E: Reorder Codes

Reorder Codes 145

DINAMAP®
ProCare Monitor
Operation Manual

DINAMAP® *ProCare* Monitor

DINAMAP® *ProCare* Monitor Operation Manual

This manual is for DINAMAP *ProCare* Monitors models 100, 200, 300, and 400, with or without printers.

- *ProCare* 100: BP, Pulse
- *ProCare* 200: BP, Pulse, and Temp
- *ProCare* 300: BP, Pulse, and SpO_2
- *ProCare* 400: BP, Pulse, Temp, and SpO_2

The model of the Monitor determines which parameters are in your monitor. Please refer to applicable sections.

Reissues and Updates

Changes occurring between issues are addressed through Change Information Sheets, Addendums, and replacement pages. If a Change Information Sheet does not accompany this manual, it is correct as printed.

Errors and Omissions

If errors or omissions are found in this manual, please notify:
GE Medical Systems *Information Technologies*
Technical Publications
4502 Woodland Corporate Boulevard
Tampa, FL 33614
1-877-274-8456

Part No. 2009360-001 A

The content of this document including all figures and drawings is proprietary information of GE Medical Systems *Information Technologies*, provided solely for purposes of operation, maintenance or repair, and dissemination for other purposes or copying thereof is prohibited without prior written consent by GE Medical Systems *Information Technologies*, Tampa, Florida.

Illustrations may show design models; production units may incorporate changes.

Hierarchy of Warnings and Cautions

A **general warning** is a statement that alerts the user to the possibility of injury, death, or other serious adverse reactions associated with the misuse of the device. A **warning** relates to steps in a procedure.

A **general caution** is a statement that alerts the user to the possibility of a problem with the device associated with its use or misuse. Such problems include device malfunction, device failure, damage to the device or damage to other property. A **caution** relates to steps in a procedure.

© GE Medical Systems *Information Technologies* 2002,
TAMPA, FL 33614

Printed in the U.S.A. All rights reserved.

United States
GE Medical Systems *Information Technologies*
4502 Woodland Corporate Boulevard
Tampa, FL 33614

Germany
GE Medical Systems *Information Technologies GmbH*
Munzinger Strasse 3
79111
Freiburg, Germany

Introduction

Introduction

About the DINAMAP® ProCare Monitor

The *ProCare* Monitor provides a small, portable, easy-to-use monitoring alternative for sub-acute hospital and non-hospital settings. The battery-operated Monitor offers noninvasive determination of systolic blood pressure, diastolic blood pressure, mean arterial pressure, pulse rate, oxygen saturation, and temperature. Monitors are available with or without integrated printers. *ProCare* Monitors are intended for use in various markets, from the physician's office to sub-acute triage and medical/surgical units.

- *ProCare* 100: BP, Pulse
- *ProCare* 200: BP, Pulse, and Temp
- *ProCare* 300: BP, Pulse, and SpO_2
- *ProCare* 400: BP, Pulse, Temp, and SpO_2

The model of the Monitor determines which parameters are in your monitor. Please refer to applicable sections.

Using the *ProCare* Monitor, a clinician can view, print, and recall clinical data that is derived from each parameter. The Monitor is also capable of alerting the clinician to changes in the patient's condition or when it is unable to effectively monitor the patient's condition. All of the main operations of the *ProCare* Monitor are easy-to-use and only a button-touch away. Please review the factory default settings and, where applicable, enter settings appropriate for your use.

Indications

The *ProCare* Monitor is intended to monitor one patient at a time in a clinical setting.

Contraindications

This device is not designed, sold, or intended for use except as indicated.

Federal law (U.S.A.) restricts this device to sale by or on the order of a clinician.

Warnings

- **Do not use the *ProCare* Monitor in the presence of magnetic resonance imaging (MRI) devices. There have been reports of sensors causing patient burns when operating in an MRI environment.**
- **Do not use the Monitor in the presence of flammable anesthetics.**
- **To help prevent unintended current return paths with the use of high frequency (HF) surgical equipment, ensure that the HF surgical neutral electrode is properly connected.**
- **To avoid personal injury, do not perform any servicing unless qualified to do so.**
- **WARNING: These Monitors should not be used on patients who are connected to cardiopulmonary bypass machines.**
- **If powering the Monitor from an external power adapter or converter, use only GE Medical Systems *Information Technologies*-approved power adapters and converters.**
- **The Monitor does not include any user-replaceable fuses. Refer servicing to qualified service personnel.**
- **To reduce the risk of electric shock, do not remove the cover or the back. Refer servicing to a qualified service person.**
- **If the accuracy of any determination reading is questionable, first check the patient's vital signs by alternate means and then check the *ProCare* Monitor for proper functioning.**

Cautions

- **Do not use replacement batteries other than the type supplied with the Monitor. Replacement batteries are available from GE Medical Systems - *Accessories and Supplies.***
- **The *ProCare* Monitor is designed to conform to Electromagnetic Compatibility (EMC) standard IEC 601-1-2, 1993 and will operate accurately in**

conjunction with other medical equipment which also meets this requirement. To avoid interference problems affecting the Monitor, do not use the Monitor in the presence of equipment which does not conform to these specifications.

- **Place the *ProCare* Monitor on a rigid, secure surface. Monitor must only be used with mounting hardware, poles, and stands recommended by GE Medical Systems *Information Technologies*.**
- **The weight of the accessory basket contents should not exceed 5 lb (2.7kg).**
- **Arrange the external AC/DC power converter, air hoses, and all cables carefully so they do not constitute a hazard.**
- **Verify calibration of BP parameter (temp and pulse oximeter do not require calibration). Ensure that the display is functioning properly before operating the *ProCare* Monitor.**
- **Do not immerse the Monitor in water. If the Monitor is splashed with water or becomes wet, wipe it immediately with a dry cloth.**
- **Do not gas sterilize or autoclave.**
- **The *ProCare* Monitor, when used with GE Medical Systems Information Technologies-approved applied parts and accessories, is protected against defibrillator damage.**

Note

- The electromagnetic compatibility profile of the *ProCare* Monitor may change if accessories other than those specified for use with the *ProCare* Monitor are used.

Product Compliance

The DINAMAP® *ProCare* Monitor is classified in the following categories for compliance with IEC 601-1:

- Internally powered or Class II when powered from external supply
- Transportable
- For continuous operation
- Not suitable for use in the presence of flammable anesthetics
- Not for use in the presence of an oxygen-enriched atmosphere (oxygen tent)
- Type BF applied parts
- IPX1, degree of protection against ingress of water
- Sterilization/Disinfection, see Appendix B
- Software is developed in accordance with IEC 601-1-4.
- This equipment is suitable for connection to public mains via power adaptors as defined in CISPR 11.
- The SpO_2 parameter conforms to EN 865:1997 with the exception of Clauses 36, 48, sub-clause 51.108.1and to ISO 9919.
- Defibrillation protected. When used with the recommended accessories, the Monitor is protected against the effects of defibrillator discharge. If monitoring is disrupted by the defibrillation, the Monitor will recover.

C US

DINAMAP® *PROCARE* MONITOR
CLASSIFIED WITH RESPECT TO ELECTRIC SHOCK, FIRE AND MECHANICAL AND OTHER SPECIFIED HAZARDS ONLY IN ACCORDANCE WITH CAN/CSA C22.2 NO. 601.1. ALSO EVALUATED TO IEC-601-2-30.

CE
0086

This product conforms with the essential requirements of the Medical Device Directive. Accessories without the CE mark are not guaranteed to meet the Essential Requirements of the Medical Device Directive.

Symbols

The following symbols are associated with the *ProCare* Monitor.

Note: The model of the Monitor determines which symbols appear on it.

Attention, consult accompanying documents

Silence

Alarms

+ / - Increase / decrease adjustable settings

Menu

Inflate/Stop

Cycle

History

Print

On/Off

Battery Power

External communications port connector

Charging

Defibrillator-proof type BF equipment

External DC power input

Class II equipment according to IEC 60536

Packaging label depicting the transportation and storage atmospheric pressure range of 500 to 1060 hPa.

IPX1

The DINAMAP® *ProCare* Monitor is protected against vertically falling drops of water and conforms with the IEC 529 standard at level of IPX1. No harmful effects will come of vertically falling drops of water making contact with the Monitor.

Getting Started

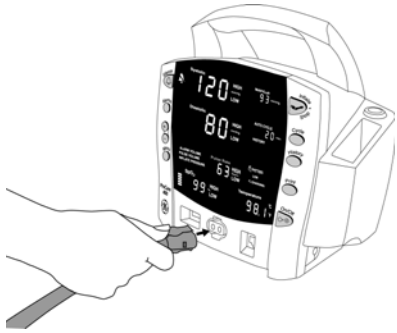
Getting Started

Unpacking the Monitor and Accessories

Before attempting to use the *ProCare* Monitor, take a few minutes to become acquainted with the Monitor and its accessories. Unpack the items carefully. This is also a good time to check for any damage or accessory shortage. If there is a problem or shortage, contact GE Medical Systems *Information Technologies*.

It is recommended that all the packaging be retained, in case the Monitor must be returned for service in the future.

Setting up BP Connections



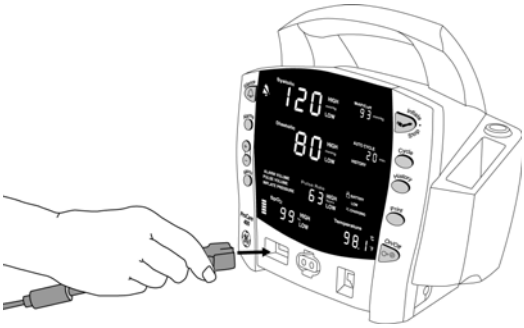
1. Connect the end of the air hose that has quick-release clips to the BP connector on the front of the Monitor. Make sure that the hose is not kinked or compressed.
   Note: To disconnect the hose from the Monitor, squeeze the quick-release clips together and pull the plug from the BP connector.
2. Select appropriate cuff size. Measure patient's limb and select appropriately sized cuff according to size marked on cuff or cuff packaging. When cuff sizes overlap for a specified circumference, choose the larger size cuff.
   Precaution: Accuracy depends on use of proper size cuff.
3. Inspect cuff for damage. Replace cuff when aging, tearing, or weak closure is apparent. Do not inflate cuff when unwrapped.
   Precaution: Do not use cuff if damaged.
4. Connect the cuff to the air hose. Refer to the *BP* section for complete cuff connection instructions.
   Warning: It is mandatory that the appropriate hose and cuff combination be used. Any attempt to modify the hose will inhibit the Monitor from switching between the neonatal and adult measurement modes.

Note: Care should be taken in reconnecting the cuff to a hose, ensuring that threads of the cuff and hose are in alignment and no cross-threading occurs.

5. Refer to the *BP* section of this manual for complete instructions on taking an accurate BP determination. **Note: Use only CRITIKON® Blood Pressure Cuffs. The size, shape, and bladder characteristics can affect the performance of the instrument. Inaccurate readings may occur unless CRITIKON® Blood Pressure Cuffs are used. Refer to Appendix E for reorder codes.**

Setting up SpO2 Connections

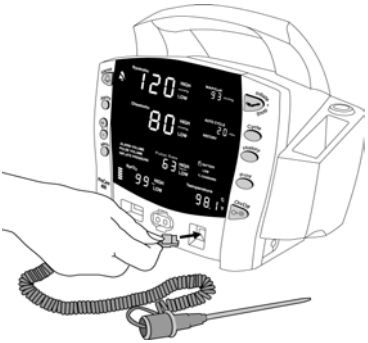
1. Plug the appropriate SpO_2 sensor into the SpO_2 sensor extension cable.
2. Then plug the SpO_2 sensor extension cable into the SpO_2 sensor connector on the Monitor.



3. Refer to the applicable *SpO_2* section of this manual for complete instructions on monitoring SpO_2.

Setting up Temperature Connections

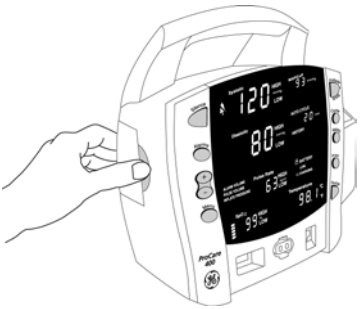
1. Connect the temperature probe cable to the temperature probe connector on the Monitor.
2. Insert the temperature probe into the probe holster at the side of the Monitor.

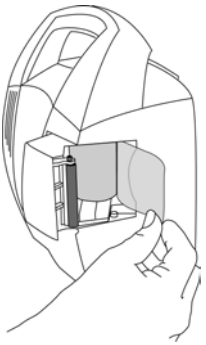


3. Refer to the *TURBO ★TEMP™* section of this manual for complete instructions on taking a temperature reading.

Setting up the Printer (Installing the Paper)

1. With the Monitor powered on, turn it so that the side with the printer is facing you.
2. While grasping the side of the Monitor, lift the printer door open by placing your thumb in the indented area and pulling. The printer door will pop open.





3.Place the roll of paper into the compartment so that the end of the paper comes off the right-side of the roll (paper is wound around the roll clockwise). Push the roll all the way to the back of the printer cavity, making sure the paper extends out of the printer cavity at least two inches.
4.Firmly press the door to close it.
5.Refer to the *Printer* section of this manual for complete instructions.

Power Sources

The *ProCare* Monitor is designed to operate from an internal lead-acid battery (see "Specifications" in *Product Overview* section).
Note: The *ProCare* Monitor is not designed to operate without an internal battery.

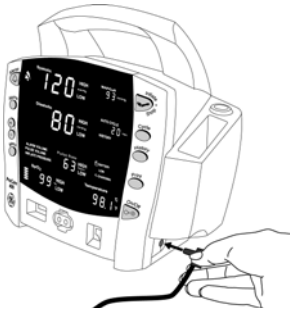
Powering the Monitor: Battery and DC Supply

Before the *ProCare* Monitor is used for the first time, the battery should be charged in the Monitor for at least 8 hours.

With external DC power connected, the green **CHARGING** indicator will light to indicate that the battery is charging.

When the Monitor is operating on battery power and the **BATTERY LOW** alarm is not active, the **BATTERY** indicator is backlit green. When the Monitor is operating on battery power and the **BATTERY LOW** alarm is active, the **BATTERY** indicator flashes green, the **LOW** indicator flashes amber, and the medium priority alarm sounds until it is acknowledged. Press the Silence button to acknowledge this alarm. Once it is acknowledged, the indicators continue to flash, but the audible alarm is silenced for 10 minutes. Once the **BATTERY LOW** condition becomes active, the Monitor should be connected to a DC power supply to recharge the battery (refer to "Power Requirements" in *Product Overview* section). If the Monitor continues to be used (approx. 10 min) without charging the battery, the Monitor enters a fail-safe mode: all functions are shutdown except for the alarm error code in the systolic window, and a constant alarm tone sounds for up to 10 minutes. After 10 minutes the alarm tone automatically shuts off and the alarm error code remains in the systolic window.

Battery charging will take place as long as the Monitor remains connected to an external DC power source.

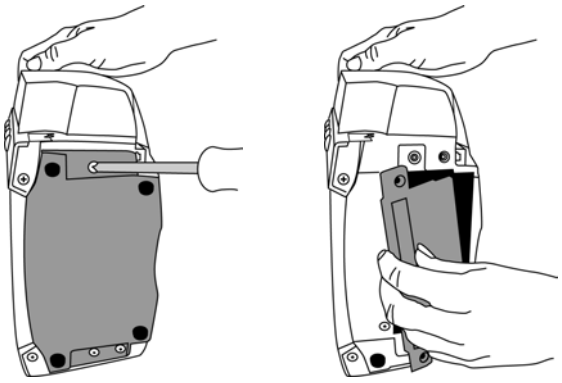


Notes

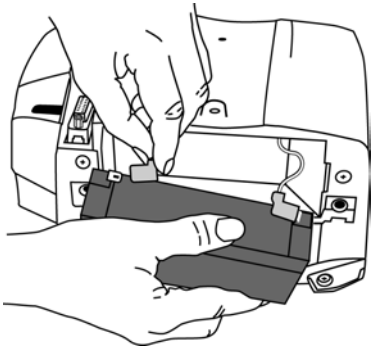
- To prolong the life of the battery, keep the Monitor connected to a DC power supply whenever possible. NEVER allow the battery to become completely discharged. A fully charged battery will power the Monitor for approximately 2 hours. To ensure full charge cycles, replace only with a recommended battery. If the Monitor is to be stored for some time, first charge the battery and then remove it and store it separately from the Monitor.

Replacing the Battery

1. Unplug the Monitor from the DC power source.
2. Looking at the bottom of the *ProCare* Monitor, remove the battery compartment cover by removing the four screws that secure the cover and help card tray.
3. Remove the help card tray and battery door cover.



3. Remove the old battery and disconnect the wires. Attach the battery wires to the new battery, ensuring the red terminal (+) is connected to the red wire and the black terminal (-) is connected to the black wire.



4. Insert the battery into the compartment.
5. Then replace the cover, help card tray, and screws. Insert the external DC power converter plug into the mains external power socket and plug into an AC outlet.

General Caution

- **Do not touch either the pin of the DC input connector or the terminals within the battery compartment and the patient at the same time.**

Switching the Monitor On and Off

To switch the *ProCare* Monitor on, push the power **On/Off** button.

As the Monitor powers up, it will run a short self-test routine, which will flash all the indicator lights and then beep the warning speaker.

To switch the Monitor off, push the power **On/Off** button again. This will terminate any measurements that may be in progress and automatically deflate the cuff. The Monitor also has an auto turn-off feature, which turns it off after 35 minutes of inactivity.

Setting the Date and Time

To set the date and time on the *ProCare* Monitor, you must first access the configuration mode. The following list shows which settings appear in which windows.

Setting:	Window	Text in Window
• Day:	Diastolic	dAY
• Month:	MAP/Cuff	MtH
• Year:	Systolic	Yr
• Hour:	min	Hr
• Minute:	min	Min

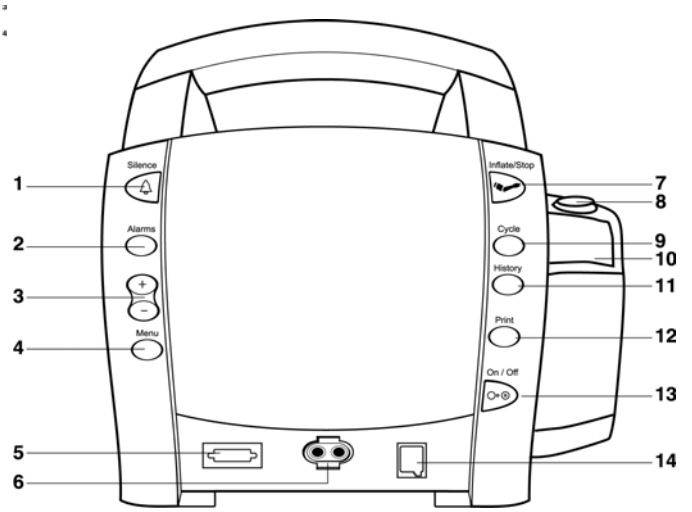
Procedures

1. With the Monitor off, press and hold the **Menu** button at the same time as pressing the **On/Off** button for 3 seconds.
2. The Monitor automatically switches on in configuration mode.

3. To set the date and time, press the **Menu** button to move from one setting to another. (You will need to press the **Menu** button several times until the **year** setting appears.) Use the **+/-** buttons to increment or decrement the individual settings. Once you are finished changing a setting, press the **Menu** button again to move to the next setting.
   Note: For the date and time to be saved, you must advance the menu through the **min** setting.
4. To exit config mode, press the **On/Off** button.

Product Overview

Product Overview



Buttons

1 Silence button: Press to mute audible alarms. Any alarm active that is acknowledgeable is also removed whenever this key is pressed. When pressed after alarm sounds (silence active), the silence icon (bell) lights to indicate that audible alarms have been silenced for 2 minutes

2 Alarms button: Press to view or adjust parameter alarm settings

3 +/- button (Plus/Minus): Press the + button to increase an adjustable setting and the - button to decrease an adjustable setting. This button is active only when a user-setting mode (limit or menu) is active

4 Menu button: Press to access menu settings that can be adjusted while in clinical mode (i.e., **ALARM VOLUME, PULSE VOLUME, INFLATE PRESSURE;** refer to *Operating Modes* in this section for a description of clinical mode)

5 SpO_2 sensor connector: SpO_2 sensor extension cable attaches here

6 BP connector: BP cuff hose attaches here

7 Inflate/Stop button: Press to start a manual BP determination or stop any BP determination

8 Temperature probe holster: Temperature probe is stored here

9 Cycle button: Press to start Auto Cycle or STAT mode

10 Temperature probe cover storage: Box of probe covers is stored here

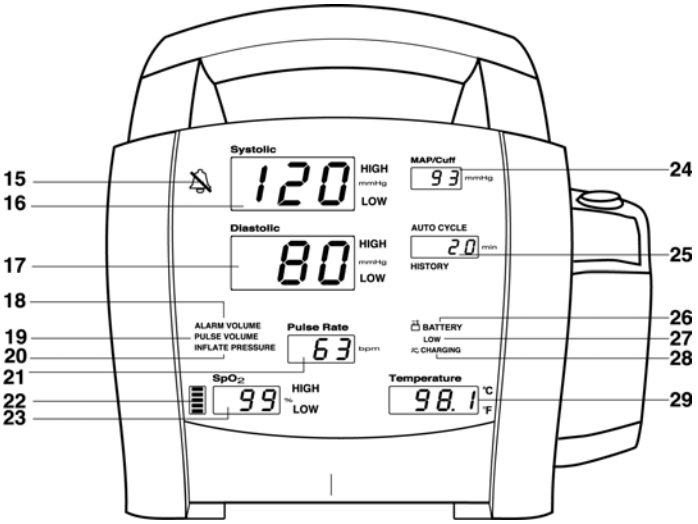
11 History button: Press to activate the history mode. When activated, it displays the most recent entries stored. Press and hold the button for 2 seconds to clear all entries stored

12 Print button: Press to print currently displayed values or all stored entries when in history mode

13 On/Off button: Controls on/off state of monitor; push for power on and push again for power off

14 Temperature probe connector: Temperature probe cable attaches here

Product Overview



Front Panel

15 Silence icon: when Silence button is pressed after alarm sounds (silence active), silence icon (bell) lights to indicate that audible alarms have been silenced for 2 minutes

16 Systolic window: 3-digit red LED indicates measured systolic BP in mmHg

17 Diastolic window: 3-digit red LED indicates measured diastolic BP in mmHg

18 Alarm volume indicator: lights to indicate you are making a change to the alarm volume

19 Pulse volume: lights to indicate you are making a change to the pulse volume

20 Inflate pressure: lights to indicate you are making a change to the inflation pressure

21 Pulse Rate window: 3-digit yellow LED shows pulse rate in beats per minute

22 SpO_2 pulse indicator: Red LED bar flashes to indicate that real-time pulse rate measurements are being derived from SpO_2 signals

23 SpO_2 window: 3-digit red LED indicates oxygen saturation in %

24 MAP/Cuff window: 3-digit red LED indicates measured MAP in mmHg and shows instantaneous cuff pressure during BP determination

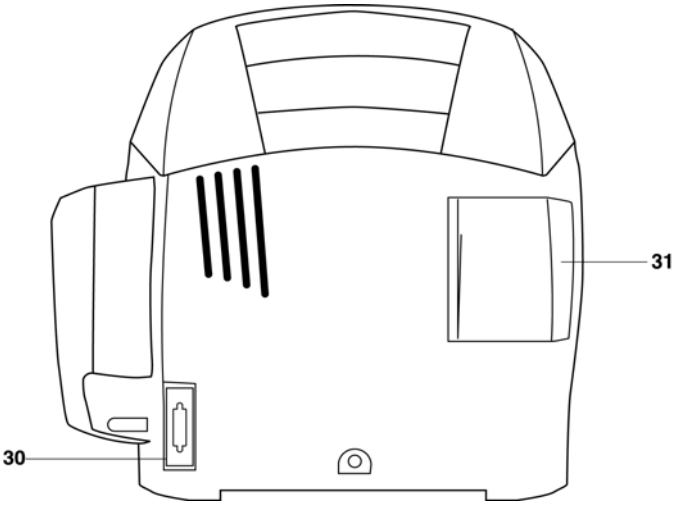
25 Min window: Displays the BP mode if manual or STAT is the cycle time when in Auto Cycle mode

26 Battery power indicator: Green LED indicates the Monitor is operating on battery power

27 Low battery power indicator: Yellow LED indicates LOW charge status of internal battery

28 Charging indicator: Green LED indicates presence of external power source and battery charging

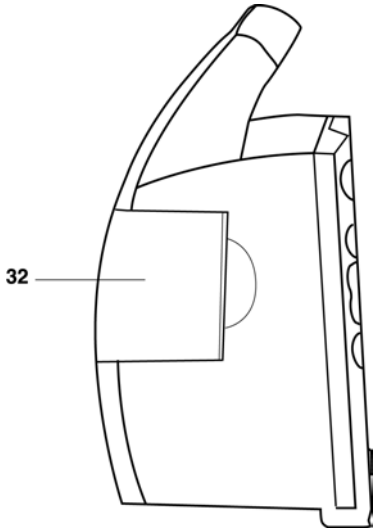
29 Temperature window: 4-digit red LED indicates measured temperature



Rear Panel

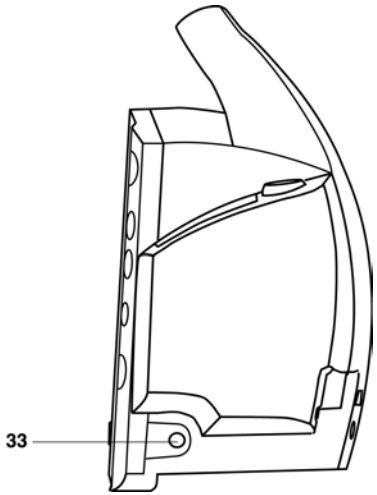
30 Data interface connector: Host communications port (15 pin D-type RS-232 serial port) for use only with equipment conforming to IEC 601-1, configured to comply with IEC 601-1-1

31 Printer compartment



Left-Side Panel

32 Printer compartment



Right-Side Panel

33 External DC power socket: To be used with approved GE Medical Systems *Information Technologies* AC-DC power converter ONLY

Windows

Each derived vital sign has an associated window for displaying the value. For each window, the vital sign's name and unit of measure are labeled above and to the right of it, respectively. An additional window--the **min** window--is available for displaying the BP mode or chosen **AUTO CYCLE** selection.

Indicators

Indicators are text messages and icons that are positioned on the front of the Monitor and can be either backlit red or green. For each vital sign that has user-adjustable limits, two indicators (**HIGH, LOW**) appear to the right of its window.

Operating Modes

The *ProCare* Monitor can operate in one of four modes: clinical, configuration, advanced configuration, and service.

Clinical mode is the Monitor's normal operating mode. While this mode is active, alarm limits and a few other commonly used settings are adjustable. All parameters are available for monitoring in this mode.

Configuration and advanced configuration modes display the software revision and allows you to configure defaults for some settings that are available in clinical mode, as well as some less commonly used settings that are only adjustable in these modes. No parameters are operable in these modes, therefore, patient monitoring should be suspended.

Entering Configuration Mode

1. With the Monitor off, press and hold the **Menu** button at the same time as pressing the **On/Off** button for 3 seconds.
2. The Monitor automatically switches on in configuration mode.

Refer to the Service Manual for instructions for use concerning advanced configuration and service mode.

Parameter Modes

The *ProCare* Monitor has three parameter modes: offline, ready, or operate.

A parameter is in offline mode when its vital sign(s) are not checked against user-set limits, alarm conditions detected by the parameter do not generate alarms, and no vital sign(s) data is displayed.

A parameter is in ready mode when its vital signs are not checked against user-set limits, alarm conditions detected by the parameter do not generate alarms, and no vital sign(s) data is displayed. However, information regarding the connection of a sensor is communicated in this mode.

A parameter is in operate mode when the appropriate vital signs are being checked against user-set limits and alarming conditions detected by the parameter generate alarms. Vital sign(s) data from the parameter is displayed.

When the Monitor is turned on and no sensor is connected, the parameter remains in offline mode. Upon detection of a sensor, the parameter auto-switches to ready mode. Upon detection of valid patient data, the parameter auto-switches to operate mode.

User Modes

The *ProCare* Monitor has four user modes that are available during clinical operating mode: **Menu, Cycle, Limit Adjustment, and History.**

The **Menu** mode allows you to access and change the three settings associated with the following indicators: **ALARM VOLUME, PULSE VOLUME, AND INFLATE PRESSURE**. To activate this mode, press the **Menu** button. With each press of the **Menu** button, the indicator appears in flashing green with the associated value appearing in red in the **Diastolic** window. As the **Menu** button is pressed, the indicators appear in the following order: **ALARM VOLUME, PULSE VOLUME, AND INFLATE PRESSURE.** To change the associated value, simply use the **+/-** button to increment or decrement, respectively. After 15 seconds of not pressing the

Menu button, the **Menu** mode is automatically exited. Otherwise, you can exit the **Menu** mode by pressing the **Menu** button one more time after viewing the oldest entry. Upon exiting **Menu** mode, the main monitoring screen is displayed. Alarm and pulse volume settings are retained after power-off. INFLATE PRESSURE is reset to its configured default after power-off.

The **Cycle** mode allows you to access Auto Cycle and STAT modes. Refer to the *BP* section for more information.

The **History** mode allows you to access the stored patient data. Refer to the *History* section for more information.

The **Limit Adjustment** mode allows you to change alarm limit settings that are used while monitoring a patient. All limit alarm settings return to their default settings after power-off. The range and increment/decrement steps for each derived vital sign that has adjustable limits are described in each parameter section. The step size specified (which cannot be adjusted) tells how much the limit value will change per increment/decrement key press and also dictates how close together a pair of limits can be.

Limit-adjustable vital signs are displayed in the following order: Systolic (HIGH, LOW), Diastolic (HIGH, LOW), Pulse Rate (HIGH, LOW), and SpO_2 (HIGH, LOW).
Note: The Temperature and MAP (mean arterial pressure) vital signs are not checked against alarm limits.

Sounds

The Monitor generates sounds based upon user interaction, parameter events, parameter and system alarms, and low battery alarms.

User interaction sounds include positive key tone and negative key tone. The positive key tone sounds one high-pitched tone when you press a button and the Monitor is able to perform the intended function. The negative key tone sounds three low-pitched tones when you press a button and the Monitor is unable to perform the intended function.

Parameter-specific sounds are related to the functioning of each parameter: NIBP, Temperature, and SpO_2. Refer to the individual parameter sections for definitions and sounds.

Alarm-specific sounds include medium and high priority. The medium-priority alarm sounds three high-pitched tones. The high-priority alarm sounds three high-pitched tones followed by two high-pitched tones. When alarms of both priorities are active, only the high-priority alarm sound is audible.

A sound is generated when AC power is lost while patient monitoring is active.

Power Sources

The *ProCare* Monitor is designed to operate from an internal lead-acid battery. For replacement rechargeable batteries, please refer to the Service section of this manual.

Product Overview

Specifications

Mechanical

Dimensions	Height: 9.7 in (24.7 cm) Width: 8.6 in (21.9 cm) without Temperature 10.0 in (25.4 cm) with Temp Depth: 5.3 in (13.5 cm)
Weight, Including Battery	5.68 lb (2.58 kg)
Mountings	Self-supporting on rubber feet
Portability	Carried by handle
Classification Information	Mode of operation: continuous Degree of protection against harmful ingress of water: Drip-proof IPX1

Power Requirements

Power Converter US:	P/N: 2009460-001 Protection against electrical shock: Class II AC input: 120 VAC/60 Hz 24W DC output voltage; 12VDC at 1A The AC mains power adapter contains a nonresettable and nonreplaceable fuse.
Power Converter UK & EUR:	P/N UK: 2008538-001 P/N EUR: 2009539-001 Protection against electrical shock: Class II AC input: 230-240 VAC/50 Hz 92mA DC output voltage; 12VDC at 1A

	The AC mains power adapter contains a nonresettable and nonreplaceable fuse.
Monitor	**Protection against electrical shock:** Internally powered or Class II when powered from specified external power supply. **DC input voltage:** 12 VDC, supplied from a source conforming to IEC 601-1. **Fuses:** The Monitor contains four fuses. The fuses are auto-resetable and mounted within the Monitor. The fuses protect the low voltage DC input, the battery, the remote alarm output, and the +5 V output on the host port connector. **Battery:** 6 volt, 3.3 amp-hours protected by internal auto-resetting fuse and thermal protection. **Minimum operation time:** 2 hrs (5 min cycle with adult cuff at 25 °C, SpO_2 active at 60 bpm, Temp in monitor mode, printout of current values every 5 min) from full charge. **Time for full recharge:** Approx. 5 hrs from full discharge when the Monitor is switched off and approx. 8 hrs when the Monitor is switched on.
Environmental	
Operating Temperature	+ 5 °C to + 40 °C (+ 41 °F to + 104 °F)
Operating Atmospheric Pressure	700 hPa to 1060 hPa

Storage Temperature	– 20 °C to + 50 °C (– 4 °F to + 122 °F)
Storage/Transportation Atmospheric Pressure	500 hPa to 1060 hPa
Humidity Range	5% to 95% noncondensing
Radio Frequency	Complies with IEC Publication 601-1-2 (April 1993) Medical Electrical Equipment, Electromagnetic Compatibility Requirements and Tests and CISPR 11 (Group 1, Class B) for radiated and conducted emissions.

IPX1

The DINAMAP® *ProCare* Monitor is protected against vertically falling drops of water and conforms with the IEC 529 standard at level of IPX1. No harmful effects will come of vertically falling drops of water making contact with the Monitor.

Printer

Printer

Description

The *ProCare* Monitor comes with an optional text-only printer. Each time a printout is started, the following information is printed: GE Medical Systems header, monitor name; model number; current software revision; a place for patient name and hand-written comments; a place for the column labels, parameter labels, or unit of measure; date; and vital signs data, if available. On some models, a pencil icon appears instead of the name and comments labels.

Installing the Paper

Refer to the *Getting Started* section for instructions.

Print Button

The **Print** button initiates two types of printouts: currently displayed values and History.

By pressing the **Print** button while the main monitoring screen is viewable, a printout of the currently displayed values is generated. Since the currently displayed values may have been derived at different times (e.g., the BP values are 5 minutes old, while Temp was just completed), the Monitor prints the value along with a time stamp of when that value was derived. Values are printed in order of the most recent to the oldest.

By pressing the **Print** button when in History mode, all entries currently stored in the history are printed in order of the most recent to the oldest.
Note: If the **Print** button was pressed during the first 10 seconds of SpO_2 monitoring, dashes will appear for SpO_2 and pulse rate readings.

The availability of the printer is determined at the time the printout is started. When the printer is unavailable, the **Print** button makes a negative key sound when you press it. The printer is unavailable if it is out of paper, too hot, or if the battery is too low, or if the Monitor is in any of the following modes: Cycle, Alarm limit adjustment, Menu, config, or service.

If no print is visible on the paper, check that the paper roll has been installed in the correct position (refer to diagram).

To tear off the printout, use a slight sideways action to pull the paper sharply up across the edge of the door.

Printer Alarms

When any of the alarm conditions occur, the alarm code flashes in the **min** window and a high-priority alarm sounds. When a printer alarm condition is active, you can acknowledge the alarm by pressing the **Silence** button.

E10: PRINTER NO PAPER
This alarm is generated when the **Print** button is pressed and the printer detects that there is no paper.

E11: PRINTER TOO HOT
This alarm is generated when a printout is in progress and the printer becomes too hot to print.

E12: BATTERY TOO LOW TO PRINT
This alarm is generated when a printout is in progress and the battery becomes too low to print.

Storage

Store thermal paper in a cool, dry place. The printed strip (thermal paper recording) should not be

- exposed to direct sunlight,
- exposed to temperatures over 100 °F/38 °C or relative humidity over 80%,
- placed in contact with adhesives, adhesive tapes, or plasticizers such as those found in all PVC page protectors.

Note: When in doubt about long-term storage conditions, store a photocopy of the thermal paper recording.

Cautions

- **The paper is thermally activated; therefore, do not store it in a hot place as discoloration may result.**
- **Use only replacement paper rolls (P/N 770137 for single rolls, 089100 for box of 10) from GE Medical Systems - *Accessories and Supplies*.**

Alarms

Alarms

The *ProCare* Monitor provides visible and audible indications of patient-, and system-related alarm conditions. An alarm can generate an audible indication, visual indication, alarm message code, and electronic record in the history.

Alarm Codes

All alarm indications are accompanied by an audible signal unless the Silence button is selected. A microprocessor system failure will generate a high-pitched audible alarm regardless of the setting of the Silence button.

There are three categories of alarms: patient-related (limit and parameter status alarms), system-related (printer, battery, and memory alarms), and system failures.

Adjusting Alarm Limits

Alarms Button

Pressing the **Alarms** button activates the limit adjustment user mode. When the limit adjustment mode is initially activated, it first displays the systolic high limit value with the **HIGH** indicator flashing in red. With each subsequent press of the **Alarms** button, the next limit in the sequence is displayed for adjustment. Press the + button to increase the displayed limit value and the **-** button to decrease the displayed limit value.

To exit limit adjustment mode after setting alarms, you must progress through all limits until you reach the last one. Then, after reaching the last limit, immediately press the **Alarms** button. The main monitoring screen appears indicating that you have exited **Alarms** mode. If monitoring is active, the current vital signs appear after you have completed your alarm setting changes. Or, you can let the mode timeout by not touching the **Alarms** button until the main monitoring screen appears indicating that you have exited **Alarms** mode. The Monitor returns to all default limit settings after power-off; these defaults are adjustable via configuration mode.

Adjustable vital signs limits are displayed in the following order: Systolic (HIGH, LOW), Diastolic (HIGH, LOW), Pulse Rate (HIGH, LOW), and SpO_2 (HIGH, LOW). Refer to each parameter section for ranges.

Note: The Temperature and MAP vital signs are not checked against alarm limits.

Adjusting the Alarm Volume

You can adjust the alarm volume by pressing the **Menu** button. **ALARM VOLUME** will flash in green with the value in the Diastolic window. You can set the **ALARM VOLUM**E range from **1** to **10** (10 being the loudest). Changing the volume applies to all alarms.

The positive key tone that sounds when you press the +/- buttons relates directly to the user-set alarm volume. Refer to *User Modes*.

Silencing and Acknowledging an Alarm

Silence Button

To silence a patient-related alarm (limit and parameter status alarms) at anytime, press the **Silence** button. The silence icon (a bell) lights to indicate that audible alarms have been silenced for 2 minutes.

Some patient- and system-related alarms are acknowledgeable. (Refer to the Alarm Message Codes table at the back of this section.) For acknowledgeable alarms that are active when the Silence button is pressed, any associated audible or visual indication is removed and any associated alarm message code is no longer displayed.

The silence icon has three states:

- Solid red: silence icon is active.
- Blinking red: alarm silence is not active and at least one alarm condition is active.
- Off: alarm silence is not active and no alarm condition is active.

Alarm Sounds

The Monitor produces two different alarm sounds based upon the priority of the alarm: medium and high priority. The medium-priority alarm sounds three high-pitched tones. The high-priority alarm sounds three high-pitched tones followed

by two high-pitched tones. When alarms of both priorities are active, only the high-priority alarm sound is audible.

Alarm Detection and Priorities

Limit Alarms

For each derived vital sign (except MAP and Temperature), the Monitor generates a limit alarm by checking its value against user-set limits. A high-limit alarm is generated when that value is greater than its high limit. A low-limit alarm is generated when that value is less than its low limit. All limit alarms are considered high priority.

Parameter	Range	Default
Systolic High	35 - 290	200
Systolic Low	30 - 285	80
Diastolic High	15 - 220	120
Diastolic Low	10 - 215	30
Pulse Rate High: NELLCOR	35 - 250	150
Pulse Rate Low: NELLCOR	30 - 245	50
Pulse Rate High: MASIMO	35 - 235	150
Pulse Rate Low: MASIMO	30 - 230	50
SpO_2 High	21 - 100	100
SpO_2 Low	20 - 99	90

When a limit violation occurs, the following happens:

- The derived vital sign that is out of limits and the associated **HIGH** or **LOW** indicator flash.
- The high priority alarm sound becomes audible unless silence is active.

Parameter Status Alarms
The Monitor generates parameter status alarms when unusual patient or sensor conditions are detected. All parameter status alarms are considered high priority alarms. Refer to the Alarm Message Codes table in this section.

When a parameter status alarm occurs, the following happens:

- Its code flashes in the associated window.
- The high priority alarm sound becomes audible unless alarm silence is active.

Printer Alarms
Refer to *Printer* section.

Memory Alarms
Memory alarms occur when battery-backed RAM is corrupted. When it occurs, all settings are reset to their factory defaults. When detected, the alarm message code appears in the **Systolic** window and the medium-priority alarm sounds.

Battery Alarms
Refer to "Powering the Monitor" in the *Getting Started* section.

System Failure Alarms
System failures are caused by a system software or hardware failure. When a system failure occurs, the error code is displayed in the **Systolic** window, and the Monitor enters a fail-safe mode in which a constant alarm tone sounds for up to 10 minutes before it is automatically switched off. Refer to the Alarm Message Codes table in this section.

Specifications

Default Settings

Alarm Volume	5

Alarm Message Code	Alarm Detected	Acknowledgeable by pressing Silence?*
E89	BP NO DETERMINATION	Yes
E85	BP LEVEL TIMEOUT	Yes
E84	BP TOTAL TIMEOUT	Yes
E83	BP PUMP TIMEOUT	Yes
E82	EXCESS AIR IN CUFF	Yes
E80	BP OVERPRESSURE	Yes
E20	SPO2 SENSOR DISCONNECTED	Yes
E21	SPO2 REPLACE SENSOR	Yes
E22	SPO2 REPOSITION SENSOR (Masimo ONLY)	Yes
E23	SPO2 SENSOR OFF FINGER	Yes
E25	SPO2 LOST PULSE	Yes
E63	TEMP DISCONNECTED	Yes
E61	TEMP PROBE BROKEN	No
E66	TEMP PROBE TOO HOT	No
E10	PRINTER NO PAPER	Yes
E11	PRINTER TOO HOT	Yes
E12	BATTERY TOO LOW TO PRINT	Yes
E00	MEMORY LOST	Yes

Alarm Message Code	Alarm Detected	Acknowledgeable by pressing Silence?*
930	SYSTEM FAILURE**	No
940	SYSTEM FAILURE**	No
950	SYSTEM FAILURE**	No
951	SYSTEM FAILURE**	No
952	SYSTEM FAILURE**	No
970	SYSTEM FAILURE**	No
971	SYSTEM FAILURE**	No
972	SYSTEM FAILURE**	No
973	SYSTEM FAILURE**	No
974	SYSTEM FAILURE**	No
975	SYSTEM FAILURE**	No

*Acknowledging an alarm by pressing the Silence button, cancels the alarm.
** Refer to the *ProCare Service Manual* for definitions and instructions.

History

History

Description

The **History** mode allows you to access the stored patient data. The *ProCare* Monitor can hold up to 25 stored entries in history. When full, the oldest entry is removed so the most recent entry can be stored. Additionally, entries are automatically removed when they become older than 24 hours.

The age of each entry is maintained and displayed in the **min** window with a minus sign (-) in front of it when other data stored for that entry is displayed. For entries that are greater than 59 minutes old, the age is displayed as HH:MM (hour:min). For entries that are less than or equal to 59 minutes old, the age is displayed in total minutes.

When viewing entries in **History** that are out of limits, the corresponding **HIGH** or **LOW** indicator appears in red.

An entry is stored in history at the completion of a BP determination and at the completion of a successful predictive temperature measurement. At the end of a BP determination, systolic, diastolic, MAP, pulse rate, and SpO_2 values are stored. At the end of a temperature determination, only the temperature value is stored.

Buttons Associated with History

To activate the **History** mode, press the **History** button. The **HISTORY** indicator flashes green while this mode is active. With each press of the **History** button, the patient data stored with the next oldest entry is displayed. Entries are displayed from the most recent to the oldest. For example, the most recent entry could have an age of -0 minutes and the oldest entry could have an age of -23:59.

After 15 seconds of not pressing the **History** button, the **History** mode is automatically exited. Otherwise, you can exit the **History** mode by pressing the **History** button one more time after viewing the oldest entry. Upon exiting **History** mode, the main monitoring screen is displayed.

Erasing Stored History

To erase stored patient data, press and hold the **History** button for a minimum of 2 seconds. All entries that were

stored in history, as well as any patient data displayed on the Monitor that relates to the previous determination or the previous temperature measurement is erased.

Windows Associated with History

Each window on the Monitor can be active during History mode. When the History button is pressed the patient data stored for each entry is displayed in the applicable windows. Patient data is displayed from most recent to oldest, indicated by the age in the **min** window.

Indicators Associated with History

The History indicator is used to show the state of the history mode. When History mode is active, the History indicator flashes green.

BP

BP

Description

The BP parameter in the *ProCare* Monitor is available with two types of BP technologies: one calibrated to intra-arterial pressure and one calibrated to the auscultatory method (specific technologies are available in select markets). Please refer to the front of your *ProCare* Monitor to see which BP technology your monitor contains. The monitors containing auscultatory technology will have an auscultatory label above the model number on the front of the Monitor. Then, refer to the *Appendix A: Principles of Nonivasive Blood Pressure* for specific information regarding both technologies. All user interface options, instructions for use, and alarms will be the same for both technologies. The BP parameter is included in all models. Blood pressure is monitored noninvasively in the *ProCare* Monitor by oscillometric method. **Note:** For neonatal populations, the reference is always the intra-arterial pressure monitoring method.

The *ProCare* Monitor has three BP modes: 1. Manual, 2. Auto Cycle, and 3. STAT. The mode is selected by the user. The actual BP determination is automated and, once it is complete, the values for systolic pressure, diastolic pressure, mean arterial pressure, and pulse rate are shown in their respective windows.

Before each BP determination, the Monitor performs a test to ensure that the cuff pressure is below a specified level. The determination is delayed until this condition is met. The Monitor senses the type of hose being used and automatically uses adult/pediatric monitoring parameters or neonatal monitoring parameters, as appropriate.

Audible and visible alarms occur when any of the values for systolic pressure, diastolic pressure, or pulse rate are outside their selected high or low limits.

Instructions for cleaning and disinfecting BP cuffs are in *Appendix B*.
Note: The Monitor will continue to operate until the battery is completely depleted in order to obtain the full use of the battery. However, if the battery reaches its “empty” point during a BP determination, it will simply stop in the middle of the determination.

General Warnings

- **Connect cuffs and inflation systems only to systems designed for non-invasive blood pressure monitoring. Devices with luers and locking luer connectors may be inadvertently connected to intravascular fluid systems that may allow air to be pumped into a blood vessel.**
- **The *ProCare* Monitor will not measure blood pressure effectively on patients who are experiencing seizures or tremors.**
- **Arrhythmias will increase the time required by the *ProCare* Monitor to determine a blood pressure and may extend the time beyond the capabilities of the Monitor.**
- **In Manual mode, the *ProCare* Monitor displays the results of the last blood pressure determination for 30 minutes or until another determination is completed. If a patient's condition changes between one determination and the next, the Monitor will not detect the change or indicate an alarm condition.**
- **Devices that exert pressure on tissue have been associated with purpura, skin avulsion, compartmental syndrome, ischemia and/or neuropathy. To minimize these potential problems, especially when monitoring at frequent intervals or over extended periods of time, make sure the cuff is applied appropriately and examine the cuff site and the limb distal to the cuff regularly for signs of impeded blood flow.**
- **Do not apply external pressure against cuff while monitoring. Doing so may cause inaccurate blood pressure values.**
- **Use care when placing cuff on extremity used to monitor other patient parameters.**
- **The *ProCare* Monitor is designed for use only with CRITIKON® dual-tube cuffs.**
- **Use only accessories approved for use with DINAMAP® Monitors. Failure to use recommended accessories may result in inaccurate readings.**
- **Blood pressure cuffs should be removed from the patient when the Monitor is powered off. If the**

extremity remains cuffed under these conditions or if the interval between blood pressure determinations is prolonged, the patient's limb should be observed frequently and the cuff placement site should be rotated as needed.

General Cautions

- **Accuracy of BP measurement depends on using a cuff of the proper size. It is essential to measure the circumference of the limb and to select the proper size cuff. In addition, the air hoses are color-coded according to patient population. The gray 12- or 24-foot hose (3.66 m or 7.3 m) is required on patients who require cuff sizes from infant through thigh cuffs. The light blue 12-foot hose (3.66 m) is required for the neonatal cuff sizes #1 through #5.**
- **If it becomes necessary to move the cuff to another limb, make sure the appropriate size cuff is used.**
- **The pulse rate derived from a BP determination may differ from the heart rate derived from an EKG waveform because the *ProCare* Monitor measures actual peripheral pulses, not electrical signals or contractions from the heart. Differences may occur because electrical signals at the heart occasionally fail to produce a peripheral pulse or the patient may have poor peripheral perfusion. Also, if a patient's beat-to-beat pulse amplitude varies significantly (e.g., because of pulsus alternans, atrial fibrillation, or the use of a rapid-cycling artificial ventilator), blood pressure and pulse rate readings can be erratic, and an alternate measuring method should be used for confirmation.**

General Notes

- A patient's vital signs may vary dramatically during the use of cardiovascular agents such as those that raise or lower blood pressure or those that increase or decrease heart rate.
- Several conditions may cause the BP parameter to calculate and display only the mean arterial pressure (MAP) without systolic and diastolic readings. These

conditions include very low systolic and amplitude fluctuations, so an accurate calculation for these values can't be made (e.g., patient in shock); too small of a difference between systolic and MAP calculations in relationship to the difference between diastolic and MAP; or a leak has occurred in the *ProCare* Monitor. If only the MAP value is displayed, an alarm message code is displayed in the systolic window, while the diastolic window remains blank.

- This equipment is suitable for use in the presence of electrosurgery.

Buttons Associated with BP

The buttons associated with BP are **Inflate/Stop** and **Cycle**.

The **Inflate/Stop** button starts and stops BP determinations. When a determination is in progress, pressing this button stops the determination. While in STAT mode, pressing this button cancels STAT mode, as well a determination if in progress. When in Auto Cycle mode, pressing this button starts a determination or cancels a determination if in progress; it does not change the mode.
Note: While an E80: BP OVERPRESSURE alarm is active, all presses of this button are ignored.

The **Cycle** button initiates the Cycle mode, which is where you can choose STAT or an auto cycle time. Cycle mode selections appear in the following order: **Stat, 1, 2, 3, 4, 5, 10, 15, 20, 30, 60, 90, 120** (minutes), and **- -** (two dashes). Choose **Stat** to start STAT mode. Choose **1-120** to select the desired cycle time and start Auto Cycle mode. When you reach the desired setting, do not press the **Cycle** button again. After 2 seconds the cycle mode is deactivated. Choose the two dashes to cancel Auto Cycle mode.
Note: While an E80: BP OVERPRESSURE alarm is active, all presses of this button are ignored.

Windows Associated with BP

The windows associated with BP are **Systolic, Diastolic, MAP/Cuff, Pulse Rate,** and **min.** The **Systolic, Diastolic, MAP/Cuff,** and **Pulse Rate** (if SpO_2 is not active) windows

are automatically cleared when a new BP determination is started. In manual mode, the displayed information is also cleared when it becomes older than 30 minutes.

The **Systolic** and **Diastolic** windows display values after a determination has completed successfully. While in STAT mode, the **Systolic** window flashes the early systolic value if it is available.

The **MAP/Cuff** window displays the derived mean arterial pressure (MAP) following the completion of a successful determination. During any type of BP determination, the pressure inside the cuff appears in this window.

The **min** window displays the BP mode of operation and the age of the previous BP determination. When both types of information are present, they flash alternately in this window. When in Manual mode, two dashes (**- -**) are displayed. When in Auto Cycle mode, the chosen Cycle time is displayed (e.g., **15**) When in STAT mode, **Stat** is displayed. When displayed, the age of the previous BP determination is preceded by a minus sign (e.g., **- 5**).

Indicators Associated with BP

The indicators associated with BP are Systolic **HIGH** and **LOW**, Diastolic **HIGH** and **LOW**, **AUTO CYCLE** and **HISTORY.**

The **AUTO CYCLE** indicator appears solid green when Auto mode is on. It flashes green when changes are being made to the current BP mode (e.g., Cycle mode is active). The **HISTORY** indicator appears solid green when the age of the previous BP determination is displayed in the **min** window.

Parameter Modes

The BP parameter is in operate mode upon power-up. It does not have a ready mode.

BP Modes of Operation

The *ProCare* Monitor has three BP modes: 1. Manual, 2. Auto Cycle, and 3. STAT. The mode is selected by the user. BP determinations are automated and, upon completion, the

values for systolic pressure, diastolic pressure, mean arterial pressure, and pulse rate are shown in their respective windows.

Manual BP Determinations
Manual mode allows you to take one blood pressure determination. Manual mode is always the BP mode of operation upon power-up. A normal, uninterrupted Manual determination takes about 40 seconds. Following a determination, the cuff pressure must drop below 5 mmHg (neonate) or 15 mmHg (adult) before another determination can be started.

Manual BP determinations are started by pressing the **Inflate/Stop** button. To stop a Manual BP determination press the **Inflate/Stop** button. The values displayed in the Systolic, Diastolic, MAP, and Pulse Rate (if SpO_2 is not active) windows are cleared after 30 minutes have lapsed.

Auto Cycle Determinations
Auto Cycle mode allows you to take multiple determinations at user-defined intervals. In the Auto Cycle mode, the pressure must be below 5 mmHg (neonate) or 15 mmHg (adult) for at least 30 seconds before the next auto determination can be started.

Auto Cycle mode is started by selecting the **Cycle** button. When in Auto Cycle mode, the AUTO CYCLE indicator appears solid green. Manual determinations can be taken while in Auto Cycle mode without affecting when the next auto determination is to start. You can also change the time interval while in Auto Cycle mode.

Once the **Cycle** button is pressed, the first Auto Cycle determination is started, and the time between determinations appears in the **min** window. You can change the time interval between Auto Cycle determinations by continuing to press the **Cycle** button until you reach the desired time interval. Time interval selections appear in the following order: **Stat, 1, 2, 3, 4, 5, 10, 15, 20, 30, 60, 90, 120** (minutes), and **- -** (two dashes)**.** When you reach the desired time interval, do not press the **Cycle** button again; after 2

seconds, the chosen time interval is retained and remains in the **min** window.

Pressing the **Cycle** button when in Auto Cycle mode activates Cycle mode again with two dashes (- -) appearing in the **min** window. If you press the **Cycle** button immediately after the first press, the next time interval appears in the **min** window. If you do not press the **Cycle** button immediately after the first press, Cycle mode is deactivated. Press the **Inflate/Stop** button to stop the determination in progress without canceling the Auto Cycle mode. Choose the two dashes (**- -**) to cancel Auto Cycle mode.

If the first Auto Cycle determination results in a limit alarm, another repeat determination is taken to verify the alarm. This occurs only once.

Whenever an Auto Cycle determination results in a E89: BP NO DETERMINATION alarm, up to nine more repeat determinations are attempted in order to achieve valid values. If at any time during this repeat cycle, the BP NO DETERMINATION alarm is acknowledged by pressing the Silence button or the Inflate/Stop button, additional determinations are not 'attempted. If the repeat cycle completes all nine repeat determinations without reaching a valid value, the Monitor returns to normal Auto Cycle mode. However, an Auto Cycle mode determination must complete successfully before a repeat cycle will follow a future Auto Cycle mode determination that results in an E89: BP NO DETERMINATION alarm.

STAT BP Determinations

STAT mode allows you to take as many determinations as possible within a 5-minute time period. The Monitor will begin another determination once the pressure is below 5 mmHg for 8 seconds (neonates) or 15 mmHg for 4 seconds (adults), unless the 5-minute period has ended or STAT mode has been canceled.

Note: BP and BP-derived pulse rate alarm limits are disabled while in Stat mode.

STAT BP determinations are started by selecting the **Cycle** button. Once the **Cycle** button is pressed, choose **Stat**. The Monitor automatically begins a 5-minute period of STAT determinations.
Note: If the Monitor was previously in Auto mode, the first STAT BP determination begins after 2 seconds.

After the first STAT determination, an early systolic value flashes in the associated window. If STAT mode is started when a determination is already in progress, that determination becomes the first in the series of STAT determinations. At the end of STAT mode, the BP mode prior to entering STAT mode is resumed. To cancel STAT mode, press the **Inflate/Stop** button.

User Settings

Mode Settings

There is one mode setting associated with this parameter: Cycle. The Cycle mode is started by pressing the **Cycle** button. While the Cycle mode is active, Cycle selections are displayed in the **min** window. Cycle selections appear in the following order: **Stat, 1, 2, 3, 4, 5, 10, 15, 20, 30, 60, 90, 120, - - .**

Limit Settings

There are two limit settings associated with this parameter: **HIGH** and **LOW**. Both limit settings are available for **Systolic** and **Diastolic** windows. The range for Systolic **HIGH** is 35 to 290 mmHg; the range for Systolic **LOW** is 30 to 285 mmHg. The range for Diastolic **HIGH** is 15 to 220 mmHg; the range for Diastolic **LOW** is 10 to 215 mmHg. The settings appear in increments of 5 mmHg.

Neonate Settings

The Monitor determines if a neonate BP cuff is being used and then checks the systolic limit and adjusts it to an appropriate limit for neonates.

Menu Settings

The **INFLATE PRESSURE** menu setting is associated with the BP parameter. This option lets you adjust the target pressure that the Monitor initially pumps to for the next

determination. The settings appear in increments of 5 mmHg. The range available is 100 mmHg to 250 mmHg. If the Monitor senses a neonate BP cuff in use, it checks the systolic limit and adjusts it to an appropriate limit for neonates.

When you change the target pressure, any values from the previous determinations are cleared from their associated windows.

Manual determinations use the value chosen for **INFLATE PRESSURE** as their target inflation pressure unless a determination is started within 2 minutes of the completion of a previous determination. For a determination that is started within 2 minutes of the previous determination, the target inflation pressure is adaptive (calculates the target inflation pressure based on the previous determination). For all Auto Cycle and STAT determinations (with the exception of the first determination of the series), the target inflation pressure is always adaptive.

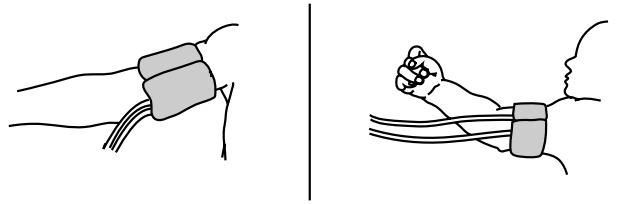
Sounds Associated with BP

There is one tone associated with this parameter: a single tone. The tone sounds at the completion of any BP determination.

Procedures

1. Connect the end of the air hose which has quick-release clips to the BP connector on the front of the Monitor. Make sure that the hose is not kinked or compressed. **Note:** To disconnect the hose from the Monitor, squeeze the quick-release clips together and pull the plug from the BP connector.
2. Select the appropriate blood pressure measurement site. Because normative values are generally based on this site and as a matter of convenience, the upper arm is preferred. When upper arm size or shape, the patient's clinical condition, or other factors prohibit use of the upper arm, the clinician must plan patient care accordingly, taking into account the patient's cardiovascular status and the effect of an alternative site

on blood pressure values, proper cuff size, and comfort. The figure shows the recommended sites for placing cuffs.
Warning: Do not place the cuff on a limb being used for intravenous infusion or any area where circulation is compromised or has the potential to be compromised.



Adult/Pediatric | Neonate

3. If patient is standing, sitting, or inclined, ensure that cuffed limb is supported to maintain cuff at level of patient's heart. If cuff is not at heart level, the difference in systolic and diastolic values due to hydrostatic effect must be considered. Add 1.80 mmHg to values for every inch (2.54 cm) above heart level. Subtract 1.80 mmHg from values for every inch (2.54 cm) below heart level.
4. Select appropriate cuff size. Measure patient's limb and select appropriately sized cuff according to size marked on cuff or cuff packaging. When cuff sizes overlap for a specified circumference, choose the larger size cuff.
   Precaution: Accuracy depends on use of proper size cuff.
   Caution: Do not use an infant cuff with an auscultatory reference DINAMAP® *ProCare* Monitor. The neonatal #5 cuff and neonatal hose may be used on patients with an arm circumference of 8 - 15 cm.
   Note: Use only CRITIKON® Blood Pressure Cuffs. The size, shape, and bladder characteristics can affect the performance of the instrument. Inaccurate readings may occur unless CRITIKON® Blood Pressure Cuffs are used.
5. Inspect cuff for damage. Replace cuff when aging, tearing, or weak closure is apparent. Do not inflate cuff when unwrapped.
   Precaution: Do not use cuff if structural integrity is suspect.
6. Connect the cuff to the air hose.
7. Inspect patient's limb prior to application.

Precaution: Do not apply cuff to areas where skin is not intact or tissue is injured.

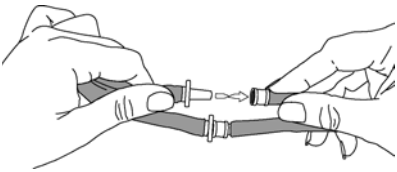
8. Palpate artery and place cuff so that patient's artery is aligned with cuff arrow marked "artery."
9. Squeeze all air from cuff and confirm that the connection is secure and unoccluded and that tubing is not kinked.
10. Wrap cuff snugly around the patient's limb. Cuff index line must fall within the range markings. Ensure that hook and loop closures are properly engaged so that pressure is evenly distributed throughout cuff. If upper arm is used, place cuff as far proximally as possible.
11. Proper cuff wrapping should be snug, but should still allow space for a finger between patient and cuff. Cuff should not be so tight as to prevent venous return between determinations.

    Warning: Using a cuff that is too tight will cause venous congestion and discoloration of the limb, but using a cuff that is too loose may result in no readings and/or inaccurate readings.
12. Proceed with monitoring in the Manual, Auto Cycle, or STAT mode.

Cuff Connections

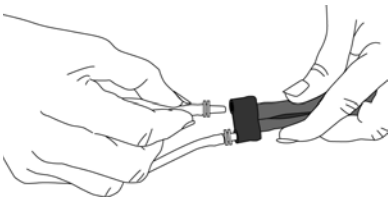
Quick-Disconnect Air Hose

Push and twist the male slip hose connector into the cuff connector to ensure a tight fit.



Neonate Air Hose

Push and twist the male slip hose connector into the cuff connector to ensure a tight fit.



Alarms

Alarm	Acknowledgeable by pressing Silence?
BP SYSTOLIC HIGH	yes
BP SYSTOLIC LOW	yes
BP DIASTOLIC HIGH	yes
BP DIASTOLIC LOW	yes
E89: BP NO DETERMINATION	yes
E85: BP LEVEL TIMEOUT	yes
E84: BP TOTAL TIMEOUT	yes
E83: BP PUMP TIMEOUT	yes
E80: BP OVERPRESSURE	yes
E82: EXCESS AIR IN CUFF	yes

The alarm codes appear in the Systolic window. When an alarm is active, it can be acknowledged by starting a new determination, with the exception of the E80: BP OVERPRESSURE alarm. BP OVERPRESSURE can only be acknowledged by pressing the **Silence** button. During STAT mode determinations, systolic, diastolic and BP-derived pulse rate values are not checked against their limits.

Critikon US Patents
4,360,029; 4,501,280; 4,546,775; 4,638,810; 5,052,397; 4,349,034; 4,543,962; 4,627,440; 4,754,761; 5,170,795

European Patents
EP122123, EP205805, EP207807

Specifications

Cuff Pressure Range (Normal operating range)	10 to 290 mmHg (adult/ped) 10 to 140 mmHg (neonate)
Blood Pressure Accuracy	Meets or exceeds ANSI/AAMI standard SP-10 (mean error ≤5 mmHg, standard deviation ≤8 mmHg)
Maximum Determination Time	120 s (adult/ped) 85 s (neonate)
Overpressure Cutoff	330 mmHg (adult/ped) 165 mmHg (neonate)
Pulse Rate Range	30 to 200 beats/min (adult/ped) 30 to 220 beats/min (neonate)
Pulse Rate Accuracy	± 3.5%

Factory Default Settings

Systolic Limits (mmHg)	HIGH: 200 LOW: 80
Diastolic Limits (mmHg)	HIGH: 120 LOW: 30
Inflation Pressure	160 mmHg (adult/ped) 110 mmHg (neonate)
Cycle button	15

All DINAMAP® Monitor regulatory and accuracy studies have been performed using CRITIKON® Blood Pressure Cuffs. Use only CRITIKON® Blood Pressure Cuffs. The size, shape, and bladder characteristics can affect the performance of the instrument. Inaccurate readings may occur unless CRITIKON® Blood Pressure Cuffs are used.

SpO_2

NELLCOR OxiMAX™ SpO2*

Description

The SpO_2 parameter in the *ProCare* Monitor is available in two different leading technologies: NELLCOR® and MASIMO SET®. Please refer to the front of your Monitor to see which SpO_2 technology your monitor contains. The SpO_2 technology logo will be on the front fascia of the Monitor. ***This section refers to NELLCOR® SpO_2 technology.***

The SpO_2 parameter is included in Models 300 and 400. To begin SpO_2monitoring, plug the sensor into the NELLCOR® DOC-10 sensor extension cable, then plug the cable into the Monitor. Simply place the SpO_2 sensor on the patient's finger; monitoring begins automatically. Functional oxygen saturation (SpO_2) of arterial blood is noninvasively and continuously monitored in the *ProCare* Monitor using pulse oximetry technology from NELLCOR®. Functional SpO_2 is the ratio of oxygenated hemoglobin to hemoglobin that is capable of transporting oxygen. This ratio, expressed as a percentage, is shown in the SpO_2 window, which is continually updated.

Pulse rate derived from SpO_2 appears in the Pulse Rate window. A tone sounds at a rate corresponding to the pulse rate and at a pitch corresponding to the SpO_2 saturation level. The pitch is highest at 100% oxygen saturation, and it continuously decreases as the saturation level falls. The Monitor displays a pulse amplitude bar. The pulse amplitude bar graph is proportional to the arterial blood flow.

Audible and visible alarms occur when SpO_2 levels are outside the alarm limits. When a parameter status alarm message code occurs, an alarm message appears in the SpO_2 window.
Note: Limit alarms, printing, and trending are not available for the first 10 seconds of SpO_2 monitoring.

*NELLCOR® is a trademark of Nellcor Puritan Bennett

General Warnings

- **Do not place SpO_2 sensor on patient during magnetic resonance imaging (MRI). Adverse reactions include potential burns to patients as a result of contact with attachments heated by the MRI radio frequency pulse, potential degradation of the magnetic resonance image, and potential reduced accuracy of SpO_2 measurements. Always remove oximetry devices and attachments from the MRI environment before scanning a patient.**
- **The use of cardio-green and other intravascular dyes at certain concentrations may affect the accuracy of the SpO_2 measurement.**
- **The SpO_2 function is calibrated to read functional arterial oxygen saturation. Significant levels of dysfunctional hemoglobins such as carboxyhemoglobin or methemoglobin may affect the accuracy of the SpO_2 measurement.**
- **Pulse oximetry readings and pulse signals can be affected by certain environmental conditions, sensor application errors, and certain patient conditions. See the appropriate sections of this manual for specific safety information.**
- **The use of accessories, transducers, and cables other than those specified may result in increased emission and/or decreased immunity and inaccurate readings of the Monitor.**

General Cautions

- **As with any clip-on sensor, pressure is exerted. The clinician should be cautious in using a clip-on sensor on patients with compromised circulation (e.g., because of peripheral vascular disease or vasoconstricting medications).**
- **Do not perform any testing or maintenance on a sensor while it is being used to monitor a patient.**
- **Bright light sources (e.g., infrared heat lamps, bilirubin lights, direct sunlight, operating room lights) may interfere with the performance of the SpO_2 function.**

To prevent such interference, cover the sensor with opaque material.

General Notes

- A patient's vital signs may vary dramatically during the use of cardiovascular agents such as those that raise or lower blood pressure or those that increase or decrease pulse rate.
- The *ProCare* Monitor that is labeled with NELLCOR® Technology is compatible only with NELLCOR® OxiMAX sensors and NELLCOR® DOC-10 sensor extension cable.
- Software development, software validation, and Risk and Hazard Analysis has been performed to a registered quality system.
- User or patient-applied parts are latex-free.

Configuration Settings Associated with SpO_2

There are two configuration settings associated with this parameter: Response Mode (**n0d**) and *SatSeconds*™ (**SAt**).

Response Mode (**n0d**) allows the user to specify the averaging technique to optimize measurements in the presence of various patient movement. Choose **Mode 1** (Normal Response) when patients are active as in exercise protocols. Choose **Mode 2** (Fast Response; default setting) for the general patient population. The configuration of this setting affects whether the pulse rate derived from SpO_2 is displayed is the Pulse Rate window while SpO_2 is in operate mode. If **Mode 1** is chosen, the pulse rate is not displayed. If **Mode 2** is chosen, the pulse rate is displayed.

With traditional alarm management, upper and lower alarm limits are set for monitoring SpO_2. During monitoring, as soon as an alarm limit is violated by as little as one percentage point, an audible alarm immediately sounds. When the % SpO_2 fluctuates near an alarm limit, the alarm sounds each time the limit is violated. To prevent these nuisance alarms, the *ProCare* Monitor uses the *SatSeconds*™ technique. The *SatSeconds*™ technique (**SAt**) limit-controls the time that the % SpO_2 level may fall outside the alarm

before an audible alarm sounds. Choose either **0, 10, 25, 50,** or **100** seconds. If **0** is chosen this limit hold-off feature is disabled.

The *SatSeconds*™ "Safety Net" is for patients with saturation levels having frequent excursions below the limit, but not staying below the limit long enough for the *SatSeconds*™ time setting to be reached. When 3 or more limit violations occur within 60 seconds, an alarm sounds even if the *SatSeconds*™ time setting has not been reached.

Buttons Associated with SpO_2

There are no buttons associated with this parameter.

Windows Associated with SpO_2

There is one window associated with this parameter: SpO_2. While in offline mode, nothing is displayed in the SpO_2 window. When in ready mode and the parameter senses that a sensor is connected, a single dash (-) appears in this window. When it is in operate mode and the parameter is reporting a valid data, the derived SpO_2 value appears in this window. The values are displayed in %.
Note: If SpO2 is the source for Pulse Rate, the Pulse Rate window is associated with this parameter.

Indicators Associated with SpO_2

There is one indicator associated with this parameter: the pulse amplitude indicator bar. The red LED bar flashes to indicate that pulse rate measurements are being derived from SpO2 signals and the height of the bar is proportional to the arterial blood flow.

Parameter Modes

The SpO_2 parameter is in offline mode upon power-up. The parameter automatically switches from offline to ready mode when a sensor is connected to the Monitor. The parameter automatically switches from ready to operate mode when pulses are detected.

If a SpO_2 SENSOR OFF FINGER or SPO2 LOST PULSE alarm is acknowledged, the parameter automatically switches from operate to ready mode. If a SpO_2 SENSOR DISCONNECTED

alarm is acknowledged, the parameter automatically switches from operate to offline mode.
Note: Once either of these alarms have been acknowledged, SpO_2 monitoring is not resumed until a valid pulse is detected again.

User Settings

Limit settings

There are two limit settings associated with this parameter: **HIGH** and **LOW**. The range for **HIGH** is 21 to 100%. The range for **LOW** is 20 to 99%. The settings appear in increments of 1%.

Menu Settings

The **PULSE VOLUME** menu setting is associated with the SpO_2 parameter. This option lets you adjust the volume of the tone that sounds after each detected heartbeat. It can be adjusted from 0 - 10 (10 being the loudest). If you set the volume to zero, no tone will sound.

Sounds Associated with SpO_2

An audible tone is provided by the Monitor for each pulse detected by the NELLCOR® SpO_2 parameter. The pitch of the audible tone is directly related to the calculated saturation value. As the saturation value increases, the pitch frequency increases continuously with each 1% increase in SpO_2. As the saturation value decreases, the pitch frequency continuously decreases. This audible tone is silenced while an alarm sounds or the **PULSE VOLUME** is set to **0**. Refer to "Menu Settings" in this section.

Procedures

1. Select a sensor that is appropriate for the patient and the clinical situation.

- **Use only NELLCOR® sensors PURPLE and WHITE and NELLCOR® DOC-10 pulse oximetery cables. Use of another pulse oximetry cable will have an adverse effect on performance. Do not attach any cable that is intended for computer use to the sensor port. Do not connect any device other than NELLCOR®-approved sensor to the sensor connector.**

Warning: Do not use a damaged sensor or one with exposed electrical contacts. Do not use sensor, cables, or connectors that appear damaged.

Note: Use only NELLCOR® OxiMAX compatible sensors, which are available from GE Medical Systems - *Accessories and Supplies*:

2. Following the directions for use supplied with the sensor, apply the sensor to the patient.

Warnings

Patient safety:

- **If the sensor is are not applied properly, the patient's skin could be injured or the ability of the *ProCare* Monitor to measure oxygen saturation could be compromised. For example, a clip-on sensor should never be taped shut. Taping the sensor could damage the patient's skin or impair the venous return, thus causing venous pulsation and inaccurate measurement of oxygen saturation.**
- **Excessive pressure from the sensor may cause necrosis of the skin.**
- **For additional warnings and information, refer to the NELLCOR® sensor's directions for use.**

Monitor performance:

- **When an SpO_2 sensor is on a limb that has a blood pressure cuff, the SpO_2 data will not be valid when the cuff is inflated. If SpO_2 readings are required during the entire blood pressure determination, attach the SpO_2 sensor to the limb opposite the one with the blood pressure cuff.**
- **Remove nail polish and artificial nails. Placing a sensor on a polished or an artificial nail may affect accuracy.**

Cautions

Patient safety:

- **Do not place any clip-on sensor in a patient's mouth or on a patient's nose or toe.**
- **Do not place a clip-on finger sensor on a patient's thumb or across a child's foot or hand.**
- **The sensor disconnect error message and associated alarm indicate that the sensor is either disconnected or**

the wiring is faulty. The user should check the sensor connection and, if necessary, replace the sensor, DOC-10 cable, or both.

- **Observe the sensor site frequently to assure adequate distal circulation. Sensor sites should be checked at least every 2 hours and rotated at least every 4 hours.**

Monitor performance:

- **Placing a sensor distal to an arterial line may interfere with adequate arterial pulsation and compromise the measurement of SpO_2.**
- **Place the sensor so that the LEDs and the photodiode are opposite each other.**

3. Plug the SpO_2 sensor into the SpO_2 sensor extension cable. Then plug the SpO_2 sensor extension cable into the SpO_2 sensor connector.
4. Proceed with monitoring. SpO_2 determinations run continuously and can run simultaneously with other measurements.

Alarms

Alarm	Acknowledgeable by pressing Silence?
SPO2 HIGH	no
SPO2 LOW	no
E20: SPO2 SENSOR DISCONNECTED	yes
E21: SPO2 REPLACE SENSOR	yes
E23: SPO2 SENSOR OFF FINGER	yes
E25: SPO2 LOST PULSE	no

Troubleshooting

This section discusses potential difficulties and suggestions for resolving them. If the difficulty persists, contact a qualified service person or your local representative.

The service manual, which is for use by qualified service personnel provides additional troubleshooting information.

PROBLEM: The pulse amplitude bar indicates a pulse, but no oxygen saturation or pulse rate values appear on the screen.
CAUSE:

- Excessive patient motion may be making it impossible for the SpO_2 function to find a pulse pattern.
- The sensor may be damaged.
- The patient's perfusion may be too low to allow the SpO_2 function to measure saturation and pulse rate.

SOLUTION:
Check the patient.

- If possible, keep the patient still; check whether the SpO_2 sensor is applied securely and properly, and replace it if necessary; move the sensor to a new site; or use a disposable adhesive sensor that may tolerate more motion.
- Replace the sensor.

PROBLEM: The SpO_2 value or the pulse rate changes rapidly; the pulse amplitude bar is erratic.
CAUSE:

- Excessive patient motion may be making it impossible for the SpO_2 function to find a pulse pattern.
- An electrosurgical unit (ESU) may be interfering with performance.

SOLUTION:
Check the patient.

- If possible, keep the patient still; check whether the sensor is applied securely and properly, and replace it if necessary; move the sensor to a new site; use a sensor that tolerates more motion.

If an ESU is interfering:

- Move the SpO_2 cable as far from the ESU as possible.
- Plug the Monitor and the ESU into different AC circuits.
- Move the ESU ground pad as close to the surgical site as possible.
- The sensor may need to be replaced with a new sensor.
- If the patient weighs less than 3 kg or more than 40 kg, apply an OXIMAX, reusable sensors (except DS-100, OXI-A/N, OXI-P/I), or OxiCliq oxygen transducer to an appropriate site. These sensors have Faraday shields which provide added protection from high electronic noise and ambient light.

PROBLEM: The oxygen saturation measurement does not correlate with the value calculated from a blood gas determination.

CAUSE:

- The SpO_2 calculation may not have correctly adjusted for the effects of pH; temperature; CO_2; fetal hemoglobin; or 2,3-DPG.
- Accuracy can be affected by incorrect sensor application or use; intravascular dyes; bright light; excessive patient movement; venous pulsations; electrosurgical interference; and placement of a sensor on an extremity that has a blood pressure cuff, arterial catheter, or intravascular line.

SOLUTION:

- Check that calculations have been corrected appropriately for the relevant variable. In general, calculated saturation values are not as reliable as direct laboratory hemoximeter measurements.
- If there is excessive light, cover the sensor with opaque material.

- Circulation distal to the sensor site should be checked routinely. Refer to the directions for use supplied with the sensor for requirements on moving the sensor to another site to ensure adhesion, skin integrity, and correct optical alignment. If skin integrity changes, move the sensor to another site.

- Try to keep the patient still, or change the sensor site to one with less motion.
- Observe all instructions, warnings, and cautions in this manual and in the directions for use of the sensor.

PROBLEM: A valid SpO_2 signal was present but has disappeared.
CAUSE:

- A BP determination on the same limb is in progress.

SOLUTION:

- An alarm message code appears on the screen, and the audible alarm will sound immediately.

PROBLEM: A REPLACE SENSOR error code has been detected.
CAUSE:

- The sensor or cable may be the wrong type or defective, the cabling may be improperly connected, or may be trying to use a wrong sensor or extension cable.

SOLUTION:
Check the patient.

- If possible, keep the patient still; check whether the proper sensor/cable is applied securely and properly, and replace it if necessary.
- Disconnect and reconnect the sensor.

PROBLEM: A SENSOR DISCONNECTED error code has been detected, but the sensor is still connected.
CAUSE:

- The sensor is not completely connected. The sensor extension cable or sensor wiring is faulty.

SOLUTION:
Check the patient.

- Check the sensor connection to the sensor extension cable and the sensor extension cable connection to the monitor. Then, if needed, replace the sensor or the sensor extension cable.

Specifications

Measurement Range

SpO_2	1 to 100%
Pulse Rate	20 to 250 beats/min

Accuracy and Motion Tolerance

Saturation	
Without Motion - Adults*	70 to 100% ±2 digits
Without Motion - Neonate*	70 to 100% ±3 digits
With Motion - Adults/Neo**	70 to 100% ±3 digits
Low Perfusion	70 to 100% ±2 digits
	0 to 69% unspecified
Pulse Rate	
Without Motion	20 to 250 beats/min ±3 digits
With Motion	normal physiologic range
	55 to 125 beats/min ±5 digits
Low Perfusion	20 to 250 beats/min ±3 digits

*Adult specifications are shown for OXIMAX MAX-A and MAX-N sensors. Neonate specifications are shown for OXIMAX MAX-N. Saturation accuracy will vary by sensor type.
**Applicability: OXIMAX MAX-A, MAX-AL, MAX-P, MAX-I, and MAX-N sensors.

NELLCOR® Sensor Accuracy

Note: All NELLCOR® sensors must be used with the NELLCOR® DOC-10 cable; the SCP-10 cable is not compatible with the *ProCare* Monitor.

Sensor Model	**SpO2 Range 70% - 100%**
OxiMAX™	
MAX-A*, MAX-AL*	±2 digits
MAX-N*†	±2 digits
MAX-P*	±2 digits
MAX-I*	±2 digits
MAX-R*‡	±3.5 digits
OxiCliq®	
OxiCliq A	±2.5 digits
OxiCliq P	±2.5 digits
OxiCliq N†	±2.5 digits
OxiCliq I	±2.5 digits

Reusable Sensor Models	
D-YS	±3 digits
D-YS with D-YSE	±3.5 digits
D-YS with D-YSPD	±3.5 digits
DS-100A	±3 digits
OXI-A/N	±3 digits
OXI-P/I	±3 digits

* The accuracy specification under motion conditions is ±3. For a definition of motion, contact NELLCOR® Technical Services or your local representative.
† The MAX-N and the OxiCliq N were tested on patients >40 kg.
‡ The accuracy specification has been determined between saturations of 80%-100%.

Neonatal Sensor Accuracy
When sensors are used on neonatal subjects as recommended, the specified accuracy range is increased by ±1 digit, to account for the theoretical effect on oximeter measurements of fetal hemoglobin in neonatal blood. For example, MAX-N accuracy on neonates is ±3 digits, rather than ±2 digits.

Sensor Light Source

Wavelength	Infrared: 890 nm (nominal) Red: 660 nm (nominal)
Power Dissipation	Infrared: 22.5 mW (max) Red: 30 mW (max)

Factory Default Settings

SpO_2 (%)	HIGH Alarm Limit: 100
SpO_2 (%)	LOW Alarm Limit: 90
Response Mode	2 (for Mode 2: Fast Response)
SatSeconds™	0

NELLCOR® Patents
US Patent No. 4,621,643; 4,653,498; 4,700,708; 4,770,179; 4,802,486; 4,869,254; 4,928,692; 4,934,372; 5,078,136; 5,351,685; 5,421,329; 5,485,847; 5,533,507; 5,577,500; 5,803,910; 5,853,364; 5,865,736; 6,083,172; Re. 35,122 and foreign equivalents.

MASIMO SET® SpO_2

Masimo Set* SpO_2
Description

The SpO_2 parameter in the *ProCare* Monitor is available in two different leading technologies: NELLCOR® and MASIMO SET®. Please refer to the front of your *ProCare* Monitor to see which SpO_2 technology your monitor contains. The SpO_2 technology logo will be on the front fascia. ***This section refers to MASIMO SET® SpO_2 technology.***

The SpO_2 parameter is included in Models 300 and 400. To begin SpO_2 monitoring. Functional oxygen saturation (SpO_2) of arterial blood is noninvasively and continuously monitored in the *ProCare* Monitor using pulse oximetry technology from MASIMO SET®. Functional SpO_2 is the ratio of oxygenated hemoglobin to hemoglobin that is capable of transporting oxygen. This ratio, expressed as a percentage, is shown in the SpO_2 window, which is continually updated.

Pulse rate when associated with SpO_2 appears in the Pulse Rate window. A tone sounds at a rate corresponding to the pulse rate and at a pitch corresponding to the SpO_2 saturation level. The pitch is highest at 100% oxygen saturation, and it becomes lower as the saturation level falls. The Monitor displays a pulse amplitude bar. The pulse amplitude bar graph is proportional to the arterial blood flow.

Audible and visible alarms occur when SpO_2 levels are outside the alarm limits. When a parameter status alarm message code occurs, an alarm message appears in the SpO_2 window.
Note: Limit alarms, printing, and trending are not available for the first 10 seconds of SpO_2 monitoring.

*MASIMO SET® is a trademark of Masimo Corporation. Possession or purchase of this device does not convey any express or implied license to use the device with replacement parts which would, alone, or in combination with this device, fall within the scope of one or more of the

General Cautions

- **If any measurement does not seem reasonable, first check the patient's vital signs by alternate means and the check the pulse oximeter for proper functioning.**
- **Inaccurate measurements may be caused by:**
  - **Incorrect sensor application or use**
  - **Significant levels of dysfunctional hemoglobins (e.g., carboxyhemoglobin or methemoglobin)**
  - **Intravascular dyes such as indocyanine green or methylene blue**
  - **Exposure to excessive illumination, such as surgical lamps (especially ones with xenon light source), bilirubin lamps, fluorescent lights, infrared heating lamps, or direct sunlight (exposure to excessive illumination can be corrected by covering the sensor with a dark or opaque material)**
  - **excessive patient movement**
  - **venous pulsations**
  - **placement of sensor on an extremity with a blood pressure cuff, arterial catheter, or intravascular line**
- **The pulse oximeter can be used during defibrillation, but the readings may be inaccurate for a short time.**
- **loss of pulse signal can occur in any of the following situation:**
  - **the sensor is too tight**
  - **there is excessive illumination from light sources such as surgical lamp, a bilirubin lamp, or sunlight**
  - **a blood pressure cuff is inflated on the same extremity as the one with a $SP0_2$ sensor attached**
  - **the patient has hypotension, severe vasoconstriction, severe anemia, or hypothermia**
  - **there is arterial occlusion proximal to the sensor**
  - **the patient is in cardiac arrest or is in shock**

MASIMO SET® SpO_2

Sensors

- **Before use, carefully read the Masimo® sensor directions for use.**
- **Use only Masimo® oximetry sensors for $SP0_2$ measurements. Other oxygen transducers (sensors) may cause improper SpO_2 performance.**
- **Tissue damage can be caused by incorrect application or use of an Masimo® sensor, for example by wrapping the sensor too tightly. Inspect the sensor site as directed in the sensor Directions for Use to ensure skin integrity and correct positioning and adhesion of the sensor.**
- **Do not use damaged Masimo® sensors. Do not use a Masimo® sensor with exposed optical components. Do not immerse the sensor in water, solvents, or cleaning solutions (the sensors and connectors are not waterproof). Do not sterilize by irradiation, steam, or ethylene oxide. See the cleaning instructions in the directions for use for reusable Masimo® sensors.**
- **Do not use damaged patient cables. Do not immerse the patient cables in water, solvents, or cleaning solutions (the patient cable connectors are not waterproof). Do not sterilize by irradiation, steam, or ethylene oxide. See the cleaning instructions in the directions for use for reusable Masimo® patient cables.**
- **Do not use the SpO_2 function during magnetic resonance imaging (MRI). Adverse reactions include potential burns to patients as a result of contact with attachments heated by the MRI radio frequency pulse, potential degradation of the magnetic resonance image, and potential reduced accuracy of SpO_2 measurements. Always remove oximetry devices and attachments from the MRI environment before scanning a patient.**
- **The use of cardio-green and other intravascular dyes at certain concentrations may affect the accuracy of the SpO_2 measurement.**
- **The SpO_2 function is calibrated to read functional arterial oxygen saturation. Significant levels of dysfunctional hemoglobins such as carboxyhemoglobin**

or methemoglobin may affect the accuracy of the SpO_2 measurement.

- **The use of accessories, transducers, and cables other than those specified may result in increased emission and/or decreased immunity and inaccurate readings of the Monitor.**
- **As with any clip-on sensor, pressure is exerted. The clinician should be cautious in using a clip-on sensor on patients with compromised circulation (e.g., because of peripheral vascular disease or vasoconstricting medications).**
- **Do not perform any testing or maintenance on a sensor while it is being used to monitor a patient.**
- **Bright light sources (e.g., infrared heat lamps, bilirubin lights, direct sunlight, operating room lights) may interfere with the performance of the SpO_2 function. To prevent such interference, cover the sensor with opaque material.**

General Notes

- A patient's vital signs may vary dramatically during the use of cardiovascular agents such as those that raise or lower blood pressure or those that increase or decrease pulse rate.
- The *ProCare* Monitor that is labeled with MASIMO SET® Technology is compatible only with MASIMO SET® sensors.
- Software development, software validation, and Risk and Hazard Analysis has been performed to a registered quality system.
- User or patient-applied parts are latex-free.

Configuration Settings Associated with SpO_2

There are three configuration settings associated with this parameter: Averaging Time (**n0d**), FastSAT **(Sat),** and Sensitivity Mode **(Sen**).

Averaging Time **(n0d)** allows you to choose the number of seconds over which SpO_2 data is averaged. Choose **4** to **16** in steps of 2.

FastSAT **(Sat)** allows you to choose **0** (for Off) or **1** (for On). If FastSAT is configured to **1** (On), the SpO_2 values are calculated quicker.

Sensitivity Mode **(Sen)** setting allows you to adjust the thresholds for calculating SpO_2 values under low perfusion conditions. Choose **1** (low perfusion-Maximized) or **2** (low perfusion-Default).

Buttons Associated with SpO_2

There are no buttons associated with this parameter.

Windows Associated with SpO_2

There is one window associated with this parameter: SpO_2. While in offline mode, nothing is displayed in the SpO_2 window. When in ready mode and the parameter senses that a sensor is connected, a single dash (-) appears in this window. When it is in operate mode and the parameter is reporting valid data, the derived SpO_2 value appears in this window. The values are displayed in %.

Note: If SpO_2 is the source for Pulse Rate, the Pulse Rate window is associated with this parameter.

Indicators Associated with SpO_2

There is one indicator associated with this parameter: the pulse amplitude indicator bar. The red LED bar flashes to indicate that pulse rate measurements are being derived from SpO_2 signals and the height of the bar is proportional to the arterial blood flow.

Parameter Modes

The SpO_2 parameter is in offline mode upon power-up. The parameter automatically switches from offline to ready mode when a sensor is connected to the Monitor. The parameter automatically switches from ready to operate mode when pulses are detected.

If a SpO_2 SENSOR OFF FINGER or SPO_2 LOST PULSE alarm is acknowledged, the parameter automatically switches from operate to ready mode. If a SpO_2 SENSOR DISCONNECTED alarm is acknowledged, the parameter automatically switches from operate to offline mode.
Note: Once either of these alarms have been acknowledged and the parameter switches to ready mode, SpO_2 monitoring is not resumed until a valid pulse is detected again.

User Settings

Limit settings

There are two limit settings associated with this parameter: **HIGH** and **LOW**. The range for **HIGH** is 21 to 100%. The range for **LOW** is 20 to 99%. The settings appear in increments of 1%.

Menu Settings

The **PULSE VOLUME** menu setting is associated with the SpO_2 parameter. This option lets you adjust the volume of the tone that sounds for each pulse detected. It can be adjusted from 0 - 10 (10 being the loudest). If you set the volume to zero, no tone will sound.

Sounds Associated with SpO_2

An audible tone is provided by the Monitor for each pulse detected by the MASIMO SET® SpO_2 parameter. The pitch of the audible tone is directly related to the calculated saturation value. As the saturation value increases, the pitch frequency increases. As the saturation value decreases, the pitch frequency decreases. This audible tone is silenced while an alarm sounds or the **PULSE VOLUME** is set to **0**. Refer to "Menu Settings" in this section.

Procedures

1. Select a sensor that is appropriate for the patient and the clinical situation.
   Warning: Do not use a damaged sensor or one with exposed electrical contacts.
   Note: Use only MASIMO® sensors, which are available from Masimo® Corporation and GE Medical Systems - *Accessories and Supplies.*

2. Following the directions for use supplied with the sensor, apply the sensor to the patient.

Warnings

Patient safety:

- **If you fail to apply the sensor properly, the patient's skin could be injured or the ability of the *ProCare* Monitor to measure oxygen saturation could be compromised. For example, a clip-on sensor should never be taped shut. Taping the sensor could damage the patient's skin or impair the venous return, thus causing venous pulsation and inaccurate measurement of oxygen saturation.**
- **Excessive pressure from the sensor may cause necrosis of the skin.**

Monitor performance:

- **When an SpO_2 sensor is on a limb that has a blood pressure cuff, the SpO_2 data will not be valid when the cuff is inflated. If SpO_2 readings are required during the entire blood pressure determination, attach the SpO_2 sensor to the limb opposite the one with the blood pressure cuff.**
- **Remove nail polish and artificial nails. Placing a sensor on a polished or an artificial nail may affect accuracy.**

Cautions

Patient safety:

- **Do not place any clip-on sensor in a patient's mouth or on a patient's nose or toe.**
- **Do not place a clip-on finger sensor on a patient's thumb or across a child's foot or hand.**
- **Observe the sensor site to assure adequate distal circulation. Sensor sites should be checked at least every 2 hours and rotated at least every 4 hours.**

Monitor performance:

- **Placing a sensor distal to an arterial line may interfere with adequate arterial pulsation and compromise the measurement of SpO_2.**
- **Place the sensor so that the LEDs and the photodiode are opposite each other.**

3. Plug the SpO_2 sensor into the SpO_2 sensor extension cable. Then plug the SpO_2 sensor extension cable into the SpO_2 sensor connector.
4. Proceed with monitoring. SpO_2 determinations run continuously and can run simultaneously with other measurements.

Alarms

Alarm	**Acknowledgeable by pressing Silence?**
SPO2 HIGH	no
SPO2 LOW	no
E20: SPO2 SENSOR DISCONNECTED	yes
E22: SPO2 REPOSITION SENSOR	yes
E23: SPO2 SENSOR OFF FINGER	yes
E25: SPO2 LOST PULSE	no

Troubleshooting

This section discusses potential difficulties and suggestions for resolving them. If the difficulty persists, contact a qualified service person or your local representative.

The service manual, which is for use by qualified service personnel provides additional troubleshooting information.

PROBLEM: The pulse amplitude bar indicates a pulse, but no oxygen saturation or pulse rate values appear on the screen.
CAUSE:

- Excessive patient motion may be making it impossible for the SpO_2 function to find a pulse pattern.
- The sensor may be damaged.
- The patient's perfusion may be too low to allow the SpO_2 function to measure saturation and pulse rate.

SOLUTION:
Check the patient.

- If possible, keep the patient still; check whether the SpO_2 sensor is applied securely and properly, and replace it if necessary; move the sensor to a new site; or use a disposable adhesive sensor that may tolerate more motion.
- Replace the sensor.

PROBLEM: The SpO_2 value or the pulse rate changes rapidly; the pulse amplitude bar is erratic.
CAUSE:

- Excessive patient motion may be making it impossible for the SpO_2 function to find a pulse pattern.
- An electrosurgical unit (ESU) may be interfering with performance.

SOLUTION:
Check the patient.

- If possible, keep the patient still; check whether the sensor is applied securely and properly, and replace it if necessary; move the sensor to a new site; use a sensor that tolerates more motion.

If an ESU is interfering:

- Move the SpO_2 cable as far from the ESU as possible.
- Plug the Monitor and the ESU into different AC circuits.
- Move the ESU ground pad as close to the surgical site as possible.
- The sensor may be damp or may need to be replaced with a new sensor.

PROBLEM: The oxygen saturation measurement does not correlate with the value calculated from a blood gas determination.

CAUSE:

- The SpO_2 calculation may not have correctly adjusted for the effects of pH; temperature; CO_2; fetal hemoglobin; or 2,3-DPG.
- Accuracy can be affected by incorrect sensor application or use; intravascular dyes; bright light; excessive patient movement; venous pulsations; electrosurgical interference; and placement of a sensor on an extremity that has a blood pressure cuff, arterial catheter, or intravascular line.

SOLUTION:

- Check that calculations have been corrected appropriately for the relevant variable. In general, calculated saturation values are not as reliable as direct laboratory co-oximeter measurements.
- If there is excessive light, cover the sensor with opaque material.
- Circulation distal to the sensor site should be checked routinely. The site must be inspected every 8 hours to ensure adhesion, skin integrity, and correct optical alignment. If skin integrity changes, move the sensor to another site.
- Try to keep the patient still, or change the sensor site to one with less motion.
- Observe all instructions, warnings, and cautions in this manual and in the directions for use of the sensor.

PROBLEM: A valid SpO_2 signal was present but has disappeared.
CAUSE:

- A BP determination on the same limb is in progress.

SOLUTION:

- An alarm message code appears on the screen, and the audible alarm will sound immediately.

PROBLEM: A REPOSITION SENSOR error code has occurred.
CAUSE:

- Weak or "noisy" signal.

SOLUTION:
Check the patient.

- If possible, keep the patient still; check whether the sensor is applied securely and properly, and replace it if necessary; move the sensor to a new site; or replace the sensor.
- Change sensor type.
- Consider increasing perfusion using heat.
- If there is excessive light, cover the sensor with opaque material.

PROBLEM: A REPLACE SENSOR error code has occurred.
CAUSE:

- The sensor or cable may be defective, or the cabling may be improperly connected.

SOLUTION:
Check the patient.

- If possible, keep the patient still; check whether the sensor/cable is applied securely and properly, and replace it if necessary.
- Disconnect and reconnect the sensor.

Specifications

Measurement Range

SpO_2	1 to 100%
Pulse Rate	25 to 240 beats/min
Perfusion Range	0.02 to 20%

Accuracy and Motion Tolerance

Saturation

Without Motion - Adult/Ped*	70 to 100% ±2 digits
Without Motion - Neonate*	70 to 100% ±3 digits
With Motion - Adult/Ped/Neo**†	70 to 100% ±3 digits
Low Perfusion‡	70 to 100% ±2 digits
	0 to 69% unspecified

Pulse Rate

Without Motion	25 to 240 beats/min ±3 digits
With Motion	normal physiologic range
	25 to 240 beats/min ±5 digits

* The Masimo SET® SpO_2 parameter with LNOP-Adt sensors has been validated for no motion accuracy in human blood studies on healthy adult volunteers in induced hypoxia studies in the range of 70-100% SpO_2 against a laboratory co-oximeter and ECG monitor. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.

**The Masimo SET® SpO_2 parameter with LNOP-Adt sensors has been validated for motion accuracy in human blood studies on healthy adult volunteers in induced hypoxia studies while performing rubbing and tapping motions at 2 to 4 Hz at an amplitude of 1 to 2 cm and a nonrepetitive motion between 1 to 5 Hz at an amplitude of 2 to 3 cm in induced hypoxia studies in the range of 70-100% SpO2 against a laboratory co-oximeter and ECG monitor. This variation equals plus or minus one standard deviation. Plus or minus one standard deviation encompasses 68% of the population.

†The Masimo SET® SpO_2 parameter with LNOP-Neo Pt sensors has been validated for neonatal motion accuracy in human blood studies on neonates while moving the neonate's foot at 2 to 4 cm against a laboratory co-oximeter and ECG monitor. This variation equals plus or minus, one standard deviation.Plus or minus one standard deviation encompasses 68% of the population.

‡The Masimo SET® SpO_2 parameter with has been validated for low perfusion accuracy in bench top testing against a Biotek Index 2 stimulator and Masimo's simulator with signal strengths of greater than 0.02% and a % transmission of greater than 5% for saturations ranging from 70 to 100%. This variation equals plus or minus, one standard deviation.Plus or minus one standard deviation encompasses 68% of the population.

Masimo® Sensor Accuracy

Sensor Model	**SpO2 Range 70% - 100%**
LNOP	
LNOP-ADT	± 2 digits
LNOP-ADT Long	± 2 digits
LNOP-PDT	± 2 digits
LNOP-NEO	± 3 digits
LNOP-NEO PT	± 3 digits
LNOP-DCI (reusable)	± 2 digits
LNOP-DCSC (reusable)	± 2 digits
LNOP-DCIP (reusable)	± 2 digits
NRI25 (reusable)	± 2 digits

Resolution	
Saturation (% SpO_2)	1%
Pulse Rate (bpm)	1

Low Perfusion Performance	
>0.02% Pulse Amplitude	Saturation (% SpO_2) ±2 digits
and % Transmission >5%	Pulse Rate ±3 digits

Interfering Substances

Carboxyhemoglobin may erroneously increase readings. The level of increase is approximately equal to the amount of carboxyhemoglobin present. Dyes, or any substance containing dyes, that change usual arterial pigmentation may cause erroneous readings.

Sensor Light Source

Wavelength	Infrared: 905 nm (nominal) Red: 660 nm (nominal)
Power Dissipation	Infrared: 22.5 mW (max) Red: 27.5 mW (max)

Factory Default Settings

SpO_2 (%)	HIGH: 100
SpO_2 (%)	LOW: 90
Sensitivity Mode	2 (for low perfusion-Default)
Averaging Time	12 seconds
FastSAT Mode	0 (for Off)

Masimo Patents

5,482,036; 5,490,505; 5,632,272; 5,685,299; 5,758,644; 5,769,785; 6,002,952; 6,036,642; 6,067,462; 6,206,830; 6,157,850, and international equivalents.

TURBO★TEMP

Description

The Temperature parameter is included in Models 200 and 400. The *ProCare* Monitor uses IVAC® TURBO★TEMP™ technology and can be used with both oral and rectal temperature probes. The TURBO★TEMP™ parameter consists of an electronic thermometer that uses a temperature-sensing device known as a thermistor. The thermistor is part of the electrical circuit and is located at the tip of the probe. To obtain temperatures, the probe tip measures the rate of change in temperature when the thermistor comes into contact with surrounding tissue. A final temperature is calculated based on this rate of change.

Temperature values are shown in the **Temperature** window in degrees Celsius or Fahrenheit, and the unit of measure is indicated by the °C °F display. The default, which is Fahrenheit, must be changed in the Configuration mode. Once in Config mode, **Unt** (abbreviation for unit) appears in the **Temperature** window; select either **°C** or **°F**.

Two modes of operation are available: predictive and monitor.

Predictive Mode

In predictive mode, a final temperature is displayed and an audible single tone sounds. Upon initiation of a measurement, the previous temperature measurement, if present, is cleared. A predictive measurement is initiated when the probe is removed from the probe holster. A predictive mode measurement is terminated when one of the following occurs:

- A final value is determined.
- The probe is inserted into the probe holster.
- The temperature measurement mode is automatically switched to monitor mode because a predictive result could not be determined.
- A Temperature alarm is issued.

A predictive temperature measurement value is automatically cleared after 2 minutes if the probe is stored

*IVAC® and TURBO★TEMP™ are trademarks of ALARIS Medical Systems

in the probe holster or 5 minutes when the probe is left out of the probe holster.

Monitor Mode
Monitor mode is most commonly used for axillary temperature determinations. In monitor mode, the display is updated continually as the patient's temperature rises or falls. Monitor mode is automatically initiated when the probe is removed from the probe holster twice within a half of a second (remove probe from holster, reinsert probe tip into holster, remove probe from holster) or when a predictive mode measurement terminates after approximately 1 minute of not being able to successfully compute a result. When in monitor mode, the temperature value flashes. When the temperature parameter switches from predictive mode to monitor mode, a tone sounds. Monitor mode is terminated when the probe is inserted into the probe holster.
Note: These temperature readings are not stored in history, printed, or reported via host comms.

General Warning

- **The performance of the Monitor may be degraded if it is operated outside of the environmental conditions specified in *Product Overview*.**

General Cautions

- **Be careful not to overextend the coiled cord of the temperature probe. Overextension can damage the probe coil connector interfaces.**
- **Accurate oral temperatures can only be obtained by placing the blue probe under the tongue in the right or left sublingual pocket. Temperatures in other locations in the mouth can vary by more than 2 °F or 1 °C.**
- **Accurate rectal temperatures can only be obtained by using the red temperature probe. Red and blue temperature probes are *not* interchangeable.**
- **Do not allow the tip of the predictive temperature probe to come into contact with a heat source (e.g., hands or fingers) prior to taking a temperature measurement. If this occurs, allow 5 seconds for the probe tip to cool before proceeding.**

- **Use only IVAC® probes and P850A probe covers. The size, shape, and thermal characteristics of the probe covers can affect the performance of the instrument. Inaccurate readings may occur unless IVAC® probes and probe covers are used.**
- **Electromagnetic Compatibility: Operating the thermometer near equipment which radiates high-energy electromagnetic and radio frequencies (electrosurgical/cauterizing equipment, portable radios, cellular telephones, etc.) may cause false alarm conditions. If this happens, reposition the thermometer away from the source of interference and perform a new measurement.**
- **If a patient's temperature is below 96.0 °F, the unit will automatically switch from the normal mode into the monitor mode within 10 seconds. It will continue to monitor the patient's temperature until the probe is removed from the patient and returned to the storage well.**

Configuration Settings Associated with Temperature

There is one configuration setting associated with this parameter: Unit of Measure (**Unt**). This setting allows you to choose °Fahrenheit (**F**) or °Celsius (**C**).

Buttons Associated with Temperature

There are no buttons associated with this parameter.

Windows Associated with Temperature

The **Temperature** window displays the value in ° C or ° F.

Indicators Associated with Temperature

The indicators associated with Temperature are **° C** or **° F**.

Measurement in Progress Indicators

Predictive Mode

• A single dash appears in the left-side of the **Temperature** window indicating use of an oral probe.

• Two dashes appear in the left-side of the **Temperature** window indicating use of a rectal probe.

• A “chase sequence” of dashes around the outside of the right-most digit of the **Temperature** window appears indicating that the probe is in contact with skin.

Monitor Mode

• The temperature value flashes indicating monitor mode.

• Four dashes flash in the **Temperature** window indicating that the measurement is < 80.0 °F.

Measurement NOT in Progress Indicators

• Two dashes appear in the center of the **Temperature** window indicating no values are present, and the probe is connected.

• Blank: The **Temperature** window appears blank indicating that no probe is connected.

Parameter Modes

The TURBO★TEMP™ parameter is in operate mode upon power-up. It does not have a ready mode.

User Settings

There are no user settings associated with this parameter.

Menu Settings

There are no menu settings associated with this parameter.

Sounds Associated with TURBO ★TEMP™

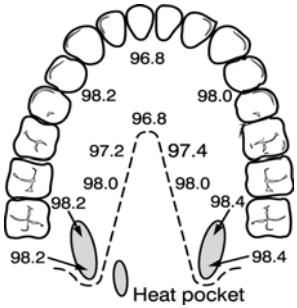
There are three sounds associated with the TURBO★TEMP™ parameter.

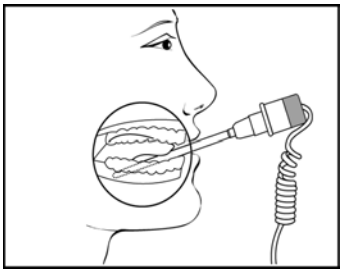
- Single tone: sounds whenever a temperature probe is removed from or inserted into the probe holster.
- Double tone: sounds whenever monitor mode is activated.
- Triple tone: sounds at the completion of a predictive temperature measurement that results in a final value.

Procedures for Oral Predictive Mode Determinations

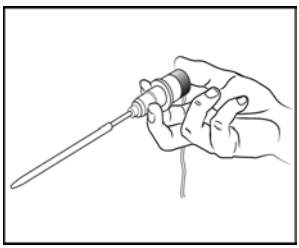
For oral temperature measurement, use the blue oral probe.

1. Connect the temperature probe cable to the temperature probe connector.
2. Holding the sides of the probe button, remove the temperature probe from the probe holster. An audible single-tone sounds. Place a protective temperature probe cover on the probe.
3. Have the patient open his/her mouth and insert the probe tip deep into the sublingual pocket where the richest blood supply is located. Failure to firmly install the probe cover may result in the probe cover becoming loose or disengaged during use. Be careful not to press the probe ejection button where the cord exits the probe as this might loosen or eject the probe cover.





4. Hold the probe steady during the entire temperature measurement process, and keep the probe tip in contact with the tissue at all times. Do not allow the patient to reposition the probe.
5. The determination begins automatically and takes approximately 10 seconds, during which time a "chase sequence" in the right-side of the **Temperature** window appears to indicate progress.
6. When the determination is complete, an audible triple tone sounds and the temperature appears on the display.
7. Remove the probe. Discard the disposable cover by holding the probe as you would a syringe and pressing the button on the probe handle. Place the probe in the probe holster. An audible single-tone sounds. Once you place the probe in the probe holster, the temperature values will be cleared in 2 minutes. They are cleared after 5 minutes when the probe is left out of the probe holster.



Procedures for Rectal Predictive Mode Determinations

For rectal temperature measurement use the red rectal probe.

1. Connect the temperature probe cable to the temperature probe connector.
2. Remove the temperature probe from the probe holster. An audible single-tone sounds. Place a protective temperature probe cover on the probe. Failure to firmly

install the probe cover may result in the probe cover becoming loose or disengaged during use. Be careful not to press the probe ejection button where the cord exits the probe as this might loosen or eject the probe cover.

3. Touch the tissue about a half inch (1.3 cm) above the sphincter muscle and carefully insert the probe, using current hospital technique for penetration. (The use of a lubricant is optional.)
4. The determination begins automatically. To ensure continuous tissue contact and maximize patient comfort, hold the probe in position until the determination is complete. This takes approximately 10 seconds, during which time a "chase sequence" in the right-side of the **Temperature** window appears to indicate progress.
5. When the determination is complete, an audible single-tone sounds and the temperature appears on the display.
6. Remove the probe. Discard the disposable cover by holding the probe as you would a syringe and pressing the button on the probe handle. Place the probe in the probe holster. An audible single-tone sounds. Once you place the probe in the probe holster, the temperature values will be cleared in 2 minutes. They are cleared after 5 minutes when the probe is left out of the probe holster.

.

Procedures for Monitor Mode Determinations (Axillary Determinations)

1. Connect the temperature probe cable to the temperature probe connector.
2. For accurate axillary measurement, the Monitor must be set in monitor mode. To trigger monitor mode, remove the probe from the probe holster, then reinsert the probe tip and remove it again within a half of a second (remove probe from holster, reinsert probe tip into holster, remove probe from holster). An audible tone sounds each time you remove the probe from the probe holster. When monitor mode is activated, an audible double-tone sounds and flashing numbers appear in the Temperature window. Place a

protective temperature probe cover on the probe and insert the probe in the patient's axilla, making sure the tip of the probe is in contact with the skin and positioned as close as possible to the axillary artery with the patient's arm held close to his/her side.

3. Leave the probe in place for the same length of time as required by standard hospital procedure for taking an axillary temperature. The Monitor does not beep to indicate a final reading.
4. Remove the probe. Discard the disposable cover by holding the probe as you would a syringe and pressing the button on the probe handle. Place the probe in the probe holster. An audible single-tone sounds. Once you place the probe in the probe holster, the temperature values will be cleared immediately.

Notes

- If there is a long delay from the time the probe is removed from the probe holster until it is inserted into the patient's mouth, it is possible that the instrument will not display a final temperature. If this occurs, insert the probe into the probe holster, remove it again, and start a new measurement.
- If an alarm is actively sounding, an audible tone will not sound.
- If the probe is not inserted/placed within 20 seconds of probe removal, Monitor mode will be entered.
- If the probe tip temperature is 92.0° F or higher (51.1° C) or 60° F (33.3° C) or lower when taken out of the probe holster, the thermometer will not be able to perform a predictive measurement. Instead, the thermometer will automatically go into monitor mode. The temperature reading will then flash. A correct final temperature reading may require 3 minutes or longer. The Monitor will not beep at final temperature. It will continue to monitor the patient's temperature until the probe is removed from the patient and returned to the probe holster.

Alarms

Alarm	**Acknowledgeable by pressing Silence?**
E63: TEMP DISCONNECTED	yes
E61: TEMP PROBE BROKEN	no
E66: TEMP PROBE TOO HOT	no

Specifications

Scale	°Fahrenheit (F) °Celsius (C)
Range	
Predictive mode	Max: 41.1° C; 106.0° F Min: 35.6° C; 96.0° F
Monitor mode	Max: 41.1° C; 106.0° F Min: 26.7° C; 80.0° F
Monitor mode accuracy	± 0.1° C ± 0.2° F (when tested in a calibrated liquid bath; meets ASTM E1112, Table 1, in range specified)
Determination time	approx. 10 seconds, typical

Use only IVAC® probes and P850A probe covers. The size, shape, and thermal characteristics of the probe covers can affect the performance of the instrument. Inaccurate readings or retention problems may occur unless IVAC® probes and probe covers are used. Refer to Appendix D for reorder codes.

Factory Default Settings

Unit of measure	° F

IVAC® Patents

U.S. D300,728, D300,909

Pulse Rate

Pulse Rate

Description

The Pulse Rate parameter is included in all Models. Pulse Rate values can be derived from one of two sources. In descending order of priority, they are pulse oximetry (SpO_2) and noninvasive blood pressure (BP). The derived values for Pulse Rate appear in the **Pulse Rate** window.

While SpO_2 is in operate mode, SpO_2 is the primary source of the Pulse Rate. At any time while SpO_2 is the source and it is unable to publish a value for Pulse Rate, three dashes (**- - -**) are displayed in **Pulse Rate** window.

BP is the secondary source of Pulse Rate. Upon completion of a BP determination, a Pulse Rate value is displayed in the **Pulse Rate** window. A Pulse Rate value is displayed as long as the results of that determination are displayed or until SpO2 switches to operate mode.
Note: When BP is in STAT mode and is the source of Pulse Rate, the Pulse Rate value is not checked against its limits upon completion of the determination.

When SpO_2 and BP are in operate mode, their associated alarms affect their availability to act as the Pulse Rate source.

General Notes

- Because the various sources measure or derive Pulse Rate differently from each other, when the Monitor changes from one source to another the value in the Pulse Rate field may change.
- A patient's vital signs may vary dramatically during the use of cardiovascular agents such as those that raise or lower blood pressure or those that increase or decrease heart rate.
- If an SpO_2-derived pulse rate is erratic, the pulse oximeter parameter may be unable to measure the pulse and may cause an alarm.

Buttons Associated with Pulse Rate

There are no buttons associated with this parameter.

Windows Associated with Pulse Rate

The **Pulse Rate** window displays the Pulse Rate value in beats per min (bpm).

Indicators Associated with Pulse Rate

The indicators associated with Pulse Rate are **HIGH** and **LOW**. Refer to "Limit Settings" in this section.

Parameter Modes

The Pulse Rate parameter is in operate mode upon power-up. It does not have a ready mode.

User Settings

Limit settings

There are two limit settings associated with this parameter: **HIGH** and **LOW**. Monitors with NELLCOR® Technology: The range for **HIGH** is 35 to 250 bpm. The range for **LOW** is 30 to 245 bpm. Monitors with MASIMO® Technology: The range for **HIGH** is 35 to 235 bpm. The range for **LOW** is 30 to 230 bpm.The settings appear in increments of 5 bpm.

Menu Settings

If SpO_2 is the source, the **PULSE VOLUME** menu setting is associated with this parameter. This option lets you adjust the volume of the pulse tones that are generated when is the SpO_2 source. It can be adjusted from 0 - 10 (10 being the loudest). If you set the volume to zero, no tone will sound.

Sounds Associated with Pulse Rate

If SpO_2 is the source, there is one sound associated with this parameter: a beat detected sound. A Pulse Rate tone is indicated by an audible beep each time a beat is detected by the SpO_2 parameter.

Note: When the pulse rate is derived from BP, there is no audible pulse beat.

Pulse Rate

Alarms

Alarm	**Acknowledgeable by pressing Silence?**
PULSE RATE HIGH	no-SpO_2, yes-BP
PULSE RATE LOW	no-SpO_2, yes-BP

SpO2 Associated Alarms (If SpO_2 is the source):

E25: SPO2 LOST PULSE
E20: SPO2 SENSOR DISCONNECTED
E21: REPLACE SENSOR

BP Associated Alarms (If BP is the source):

E89: BP NO DETERMINATION
E85: BP LEVEL TIMEOUT
E84: BP TOTAL TIMEOUT
E83: BP PUMP TIMEOUT
E80: BP OVERPRESSURE
E82: EXCESS AIR IN CUFF

Specifications

Factory Default Settings

Pulse Rate	HIGH: 150
	LOW: 50

Refer to individual SpO_2 and BP sections.

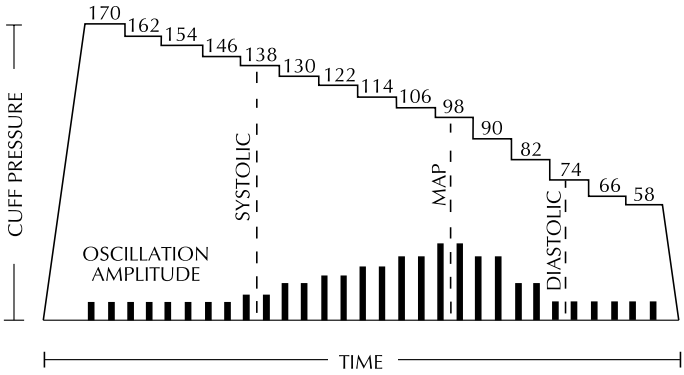
Appendix A: Principles of Noninvasive Blood Pressure

Appendix A

Principles of Noninvasive Blood Pressure Determination

The oscillometric method of determining NIBP is accomplished by a sensitive transducer, which measures cuff pressure and minute pressure oscillations within the cuff. The first determination sequence initially pumps up to a cuff pressure of about 160 mmHg for adult/pediatric patients or 110 mmHg for neonates depending on initial target pressure preset. After inflating the cuff, the Monitor begins to deflate it and measures systolic pressure, mean pressure, and diastolic pressure. When the diastolic pressure has been determined, the Monitor finishes deflating the cuff and updates the screen.

The Monitor deflates the cuff one step each time it detects two pulsations of relatively equal amplitude. The time between deflation steps depends on the frequency of these matched pulses (pulse rate of the patient). However, if the Monitor is unable to find any pulse within several seconds, it will deflate to the next step. The process of finding two matched pulses at each step provides artifact rejection due to patient movement and greatly enhances the accuracy of the Monitor. The figure shows the NIBP determination sequence.

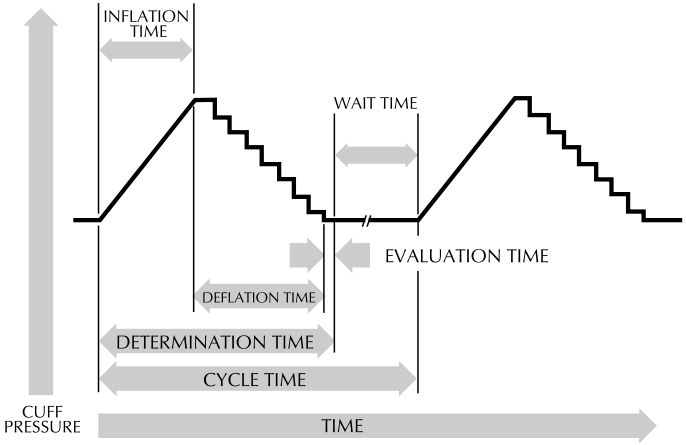


NIBP Determination Sequence

At each step the microprocessor stores cuff pressure, the matched pulse amplitude, and the time between successive pulses. The stepped deflation and matched pulse detection

continues until diastolic pressure is determined or total cuff pressure falls below 7 mmHg. The Monitor then deflates the cuff (to zero detected pressure), analyzes the stored data, and updates the screen.

The operating cycle is composed of four parts: inflation time, deflation time, evaluation time, and wait time. Wait time, which varies from mode to mode, is affected by the cycle time (auto mode) or operator intervention (manual mode). The figure shows the basic operating cycle.



NIBP Operating Cycle

Systolic Search

If systolic pressure is not found, the Monitor can search at cuff pressures higher than the initial target pressure. If the determination is in a late stage, the Monitor will inflate the cuff to 70 mmHg above the initial target pressure to get better data in the systolic region. If the determination is in an early stage, the Monitor will inflate the cuff to 50 mmHg above the initial target pressure. The maximum pressure allowed in systolic search is limited by the normal range for cuff pressures.

In any operating mode, if a patient's systolic pressure exceeds the inflation pressure of the monitor, the Monitor will begin normal deflation sequence, detect the absence of

a systolic value, stop deflation, reinflate to a higher (than initial) inflation pressure (290 mmHg maximum), and resume normal deflation sequence. This additional inflation will occur only once per determination.

If a previous valid systolic pressure is displayed and less than 2 minutes old, and the new systolic pressure oscillations are compared with the previous valid determination and the Monitor "thinks" that the systolic was not obtained, the Monitor will inflate the cuff to a pressure of an additional 50 mmHg above the immediately preceding inflation. This additional inflation will occur only once per determination.

Reference Used to Determine NIBP Accuracy
To establish accuracy of an NIBP device, manufacturers have used several different types of references. The reference blood pressures may be obtained by invasive pressure monitoring at the central aortic region or at the radial sites. The reference blood pressures may also be obtained by noninvasive methods like auscultatory method (using cuff and stethoscope).
Note: For neonatal populations, the reference is always the invasive pressure monitoring method.

DINAMAP® Monitors With Intra-Arterial Reference (DINAMAP® Classic Technology)
For these monitors, the NIBP is referenced to the invasive blood pressure obtained at the central aortic region.

DINAMAP® Monitors With Auscultatory Reference (DINAMAP® Auscultatory Technology)
In these monitors, the reference blood pressure is the auscultatory method for adult and pediatric populations. For neonatal populations, the reference is the invasive blood pressure obtained at the central aortic region.
Caution: Do not use an infant cuff with an auscultatory reference DINAMAP® *ProCare* Monitor. The neonatal #5 cuff and neonatal hose may be used on patients with an arm circumference of 8 - 15 cm.

Appendix B: Maintenance

Appendix B

Maintenance

Cleaning the Monitor

The Monitor and accessories are to be kept clean and used according to the instructions provided here and in the Service Manual.

The exterior of the Monitor may be wiped clean with a soft cloth slightly dampened with mild detergents. The Monitor and accessories should be inspected once yearly for wear and damage.

- Do ***not*** immerse unit.
- Do ***not*** clean with isopropyl alcohol or other solvents.
- Do ***not*** immerse hoses.

Cuff Cleaning and Disinfection

General

The cuff must be thoroughly cleaned with the specified detergent before reuse. The additional use of household bleach as described below provides at least intermediate-level disinfection.

- Apply cuff hose plugs before cleaning.
- The following cleansing procedure was repeated 20 times on DURA-CUF® Blood Pressure Cuffs and once on SOFT-CUF® Blood Pressure Cuffs without affecting the performance of the cuff.
- While this procedure is adequate for cleaning/ disinfection, it may not remove all stains.
- Do ***not*** immerse hoses.
- Do ***not*** immerse cuffs without prior application of cuff hose caps.

Materials

- Enzymatic detergent such as ENZOL* enzymatic detergent (US) or Cidezyme* enzymatic detergent (UK)
- Distilled water
- 10% solution of household bleach (5.25% sodium hypochlorite) in distilled water
- Soft cloths and soft-bristled brushes

*Trademark

- Spray bottles

Procedure

1. Prepare the enzymatic detergent according to the manufacturer's instructions and the 10% bleach solution, in separate spray bottles.
2. Spray the detergent liberally on device. If the material is dried on, allow the cuff to sit for 1 minute. For soil on the soft part of the closure or the cuff itself, wipe the material off with a soft cloth. For persistent contamination on the soft part of the closure, use a soft-bristled brush to loosen particles. Rinse with copious amounts of distilled water. Repeat until no visible contamination remains. For soil on the hook part of the closure, use a soft-bristled brush to remove the material, and rinse with copious amounts of distilled water. Repeat until no visible contamination remains.
3. Spray the 10% bleach solution on the affected area until the area is saturated. Allow the cuff to sit for 5 minutes.
4. Wipe away any excess solution and rinse the cuff again with distilled water. Allow 2 hours for drying.

The user has the responsibility to validate any deviations from the recommended method of cleaning and disinfection.

For additional information on infection control procedures, contact GE Medical Systems *Information Technologies* Technical Support.

Temperature Devices

Do not immerse predictive temperature probes. The probe may be cleaned with a solution of 10% bleach in water. Use a cloth or sponge—just damp, not wet—and avoid getting any liquid into the interior of the probe.

SpO_2 Sensors

Adhesive sensors are sterile and for single use only. Reusable sensors should be cleaned before reuse with a 70% alcohol solution. If low-level disinfection is required, use a 1:10 bleach solution. Do not use undiluted bleach (5% - 5.25% sodium chlorite) or any cleaning solution other than those recommended here because permanent damage to the

sensor could occur. Do not sterilize the sensor by irradiation, steam, or ethylene oxide. If disposable sensors or their packaging are damaged, they must be disposed of as advised in this appendix.

To clean or disinfect the sensor:

1. Saturate a clean, dry gauze pad with the cleaning solution. Wipe all surfaces of the sensor and cable with this gauze pad.
2. Saturate another clean, dry gauze pad with sterile or distilled water. Wipe all surfaces of the sensor and cable with this gauze pad.
3. Dry the sensor and cable by wiping all surfaces with a clean, dry gauze pad.

Storage and Battery Care

If it becomes necessary to store the Monitor for an extended period of time, first fully charge then remove the battery. Then store the Monitor and the battery in the original packaging materials.

Batteries should always be fully charged before being placed in storage. Even after 6 months of storage, a fully charged battery can retain about 80% of its charge. A fully charged battery in good condition will provide sufficient power to operate a Monitor for approximately 2 hours, including temperature and BP measurements made at 5-minute intervals.

It is best to keep the battery charged as fully as practical and never store the Monitor with the battery in a discharged condition. When the battery will no longer hold a charge, remove and replace it with one of the same part number. Failure to replace the battery with the same GE Medical Systems *Information Technologies* part number may result in shorter battery life.

Battery charging will take place as long as the Monitor remains connected to an external DC power source.

Cautions

- **To ensure that the battery will be ready for portable operation, keep the Monitor connected to a mains supply whenever possible.**
- **Repeated failure to fully charge the battery will result in a significant reduction in battery life.**
- **The expected lifetime of the battery largely depends on the way in which the Monitor is used. If the battery is allowed to completely discharge before being fully recharged, the battery should survive around 700 charge/discharge cycles.**

Replacement batteries may be obtained from GE Medical Systems *Information Technologies*.
Note: The replacement part number of the battery is 633178CR. Do not use other types.

Fuses

The Monitor contains four fuses. The fuses are auto-resetable and mounted within the Monitor. The fuses protect the low voltage DC input, the battery, the remote alarm output, and the +5 V output on the host port connector.

Calibration

Calibration of the BP parameter should be checked at least once a year or when there is doubt about the validity of the readings.

Leak Testing

A leak test of the BP parameter should be performed at least once a year or when there is doubt about the validity of the pressure readings.

Caution: Refer calibration and leak testing to qualified service personnel. Full calibration details are available in the DINAMAP® *ProCare* Monitor Service Manual, available from GE Medical Systems *Information Technologies*.

Appendix B

Disposal of Product Waste

As you use the *ProCare* Monitor, you will accumulate solid wastes that require proper disposal or recycling. These include batteries, patient applied parts, and packaging material.

Batteries

Caution: Do not incinerate batteries.

The sealed rechargeable battery contains lead and must be recycled. Discharge this battery prior to disposal. Place the battery in packaging which electrically isolates its contents. Do not puncture or place the battery in a trash compactor. Do not incinerate the battery or expose it to fire or high temperatures. Recycle in accordance with applicable regional body controlled guideline.

In the USA, call 1.800.SAV.LEAD for instructions on how to recycle. For international users, contact your local authorities on recycling. GE Medical Systems *Information Technologies* encourages the recycling of lead-acid batteries.

Patient Applied Parts

Certain patient applied parts, such as those with adhesive (disposable SpO_2 sensors), are intended for single use and should be disposed of properly as medical waste in accordance with regional body controlled guideline.

Other patient applied parts, such as blood pressure cuffs, should be cleaned according to instructions. Inspect reusable applied parts for wear, replace as necessary, and dispose of used product as medical waste in accordance with regional body controlled guideline.

Packaging Material

Retain original packaging materials for future use in storing or shipping the Monitor and accessories. This recommendation includes corrugated shippers and inserts.

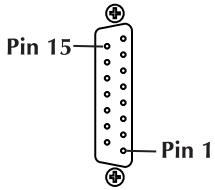
Whenever possible recycle the packaging of accessories and patient applied parts.

Appendix C: Connection Details

Appendix C

Connection Details

Host Port Connector (rear panel)



WARNING! Auxiliary equipment connected to the DINAMAP® *ProCare* Monitor will result in the formation of an electromedical system and thus, must comply with the requirements of EN 60601-1-1/ IEC 601-1. All host port signals are NON-ISOLATED and should be connected to equipment conforming to IEC-601-1, configured to comply with IEC 601-1-1 ONLY. Where isolation of data communication is required, GE Medical Systems *Information Technologies* part number ILC232 should be used. If external alarm control is required, GE Medical Systems *Information Technologies* part number 487208 (Isolated Remote Alarm Cable Assembly) should ALWAYS be used. Please refer to the Information Sheet included with the isolated remote alarm cable for operational details.

Note: When using remote alarm, the *ProCare* Monitor should be considered the primary alarm source. The secondary alarm is used for secondary purposes only.

Pin #	Function
1	Common
2	Inverted TTL Transmit Data
3	Inverted TTL Receive Data
4	+5 volts
5	No connection
6	No connection
7	Common
8	Remote Alarm
9	No connection
10	No connection
11	RS232 Transmit Data (TxD)
12	No connection
13	RS232 Receive Data (RxD)
14	No connection
15	No connection

Appendix D: Warranty, Service, & Spare Parts

Appendix D

Warranty, Service, and Spare Parts

Warning: There are no user serviceable parts inside the DINAMAP® *ProCare* Monitor. Refer all servicing to qualified personnel.

Warranty

All repairs on products under warranty must be performed or approved by GE Medical Systems *Information Technologies*. Unauthorized repairs will void the warranty.

Products not covered by warranty should be repaired only by qualified electronics service personnel.

Assistance and Parts

If the product malfunctions or if assistance, service, or spare parts are required, contact GE Medical Systems *Information Technologies* Technical Support.

Before contacting GE Medical Systems *Information Technologies* it is helpful if you can duplicate the problem and check and confirm the operation of all accessories to ensure that they are not the cause of the problem.

When calling, please have the following information at hand:

- product name and model number and complete description of the problem
- the serial number of your Monitor
- your name and address
- a purchase order number if out-of-warranty repairs or spare parts are required
- your GE Medical Systems *Information Technologies* account number, if applicable
- the part number for spare or replacement parts

Repairs

If your product requires warranty, extended warranty, or non-warranty repair service, call GE Medical Systems *Information Technologies* and a representative will assist you.

To facilitate prompt service in cases where the product has external chassis or case damage, please advise the representative when you call.

The representative will record all necessary information and will provide a Return Authorization Number. Prior to returning any product for repair, a Return Authorization Number must be obtained.

Packing Instructions

If you have to return goods for service, follow these recommended packing instructions:

- Remove all hoses, cables, sensors, power cords, and ancillary products from the Monitor before packing, unless you suspect they are associated with the problem.
- Wherever possible use the original shipping carton and packing materials.
- Observe the environmental conditions detailed in *Product Overview*.

It is recommended that all returned goods be insured. Claims for loss or damage to the product must be initiated by the sender.

Service Manuals

Service Manuals containing calibration and repair information can be ordered from GE Medical Systems *Information Technologies*. These manuals also include assembly drawings, and spare parts lists. The reorder number of the Service Manual is 2009381-001.

Appendix E: Reorder Codes

Appendix E

Reorder Codes

PRODUCT	PROD CODE	ECAT
DINAMAP *ProCare* Monitor Operations Manual-English	2009360-001	
DINAMAP *ProCare* Monitor Service Manual	2009381-001	
Battery	633178CR	
Power Convertor US	2009460-001	
Power Convertor UK	2008538-001	
Power Convertor EUR	2009539-001	
Printer Paper (box of 10)	089100	E9050KP
DINAMAP Rolling Stand	003215	E9050JB
Pole Mounting Option	2009762-001	
Wall Mounting Option	2009763-001	
Hose Management Option	2009764-001	
Bed Rail Option	2009765-001	
Nurse Call	487208CR	
NIBP:		
Air Hose 12 ft Adult/Pediatric, Screw Connector	107365	E9050LH
Air Hose 24 ft Adult/Pediatric, Screw Connector	107366	E9050LJ
Air Hose 12 ft Neonatal, Quick Disconnect	107368	E9050LK
Air Hose 12 ft Adult/Pediatric, Quick Disconnect	88847	E9050KN
CUFF Assortment Packs:		
CLASSIC-CUF® Assortment Pack Includes: 1 each: Infant, Child, Small Adult, Adult, Large Adult, Thigh Cuff	2692	E2692J
CLASSIC-CUF® Assortment Pack, Neonate Includes: 2 Neo #1, 3 Neo #2, 5 Neo #3, 5 Neo #4, 5 Neo #5	2693	E2693J
SOFT-CUF® Assortment Packs: Includes: 1 Infant, 1 Child, 2 Small Adult, 2 Adult, 2 Large Adult, 1 Thigh, 1 Adult Long Cuff	2695	E2695J
SOFT-CUF® Assortment, Neonate Includes: 2 Neo #1, 3 Neo #2, 5 Neo #3, 5 Neo #4, 5 Neo #5	2694	E2694J
DURA-CUF® Assortment Packs: Includes: 1 each: Infant, Child, Small Adult, Adult, Large Adult, Thigh Cuff	2699	E2699J
DURA-CUF® Assortment Pack, Adult Includes: 1 each: Infant, Child, Small Adult, Adult, Large Adult, Thigh Cuff	2698	E2698J
DURA-CUF® Assortment Pack, Child Includes: 2 Infant, 3 Child, and 1 Small Adult Cuff	2697	E2697J

Additional Blood Pressure Cuff Codes are available through: www.gemedicalsystems.com

TEMPERATURE:	
IVAC* Turbo Temp Oral Temperature Probe, Long Cord	2008774-001
IVAC* Turbo Temp Rectal Temperature Probe, Long Cord	2008775-001

*IVAC is a trademark of Alaris Medical Systems

IVAC* Temperature Probe Covers	88015	E9050KK

SPO2:

NELLCOR:*

Pulse Oximeter Cable DOC-10	2008773-001
DuraSensor Adult Oxygen Sensor	DS100A

*Masimo**:*

Adult Reusable Sensor, 1/Bx (NR125)	2009745-001
Cable (PC08)	2009743-001

*NELLCOR is a trademark of Nellcor Puritan Bennett.
**Masimo is a trademark of Masimo Corporation

CE

2009360-001 A